

Keynote address from TÜV SÜD and Amgen at Medical Devices and IVD Conference in November!

SMi Group reports: Navigating new legislations and strategies to comply with regulatory requirements

LONDON, LONDON BRIDGE , UNITED KINGDOM, July 19, 2021

[/EINPresswire.com/](https://www.einpresswire.com/) -- The MDR and IVDR updates will have a profound impact on the [medical](#) device industry, bringing certain products into regulatory scope that were previously excluded, introducing new manufacturing requirements, and increasing the burden for post-market surveillance. What does this mean for the future of the Medical [Devices](#) and [IVD](#) industry?



More information on the programme and speaker line-up can be found here: www.medicaldevices-ivd.com/einpr3

IVDR implementation: State of play, the Notified Body view

- Status of IVDR implementation: Where do we see the bottlenecks.
 - Lessons learnt from IVDR certification projects
 - What should manufacturers consider during the remaining time until DoA
- Andreas Stange, Vice President, TÜV SÜD

Combination Products: End-to-End Risk Management

- Product Development and Lifecycle Management are founded on principles of risk management
- Key considerations for Risk Management for Combination Products

Susan Needle, Executive Director, Combination Products, Devices, Diagnostics & Digital Health Regulatory Affairs, Amgen

BENEFIT OF ATTENDING:

- Engage with notified body and competent authority representatives addressing key MDR and IVDR requirements
- Gain insights from pharma regulatory experts in compliance and companion diagnostics
- Understand the latest guidance on Post-Brexit IVD Regulations to overcome common challenges, in addition to considering evolving global regulations
- Examine post-market expectations for surveillance and vigilance of your medical devices
- Delve into how COVID-19 has impacted the MDR and IVD regulations

Register online by 30th September and save £100: www.medicaldevices-ivd.com/einpr3
#MedicalDevicesIVD

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