



## **Angela Stokes**

I have almost 30 years of experience in medicinal product and medical device development, including extensive experience of ICH and CHMP guidelines and also ISO standards for medical devices and in vitro diagnostics. I have written several chapters for books on drug development and Regulatory Affairs and write for several peer-reviewed journals including the Regulatory Rapporteur. I speak on a variety of MSc courses and at global conferences on a regular basis. I was honoured with a TOPRA award for Regulatory Excellence in 2015 and am currently President of TOPRA. Currently I am a Senior Director of Global Regulatory Consulting at Syneos Health.

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