

One-Voice-of-Quality (1VQ) Solutions Part 2 - Podcast Episode



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By

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This is a follow-up to Part 1 of this series, where we introduced the concept of One-Voice-of-Quality to our listeners and had a chance to really lay out the ramifications of failure to change, not only in the Post Approval Change process, but the mindset around the implementation of a global regulatory framework. In this latest episode - Part 2, you will hear from Emma Ramnarine once again as we expand on details for a greater understanding around the how's and who's that are necessary to this One-Voice-of-Quality initiative.

Read more from Emma on this topic:

1. [Industry One-Voice-of-Quality \(1VQ\) Solutions: Effective Management of Post-Approval Changes in the Pharmaceutical Quality System \(PQS\)—through Enhanced Science and Risk-Based Approaches](#)
2. [Industry One-Voice-of Quality Concept Paper: Solving the Global Continual Improvement and Innovation Challenge: How an Effective Pharmaceutical Quality System Can Transforms Post-Approval Change Management](#)
3. [Continual Improvement While Maintaining A State of Control: A Concealed Paradox or a Mutual Interdependence?](#)
4. [Demonstrating Pharmaceutical Quality System Effectiveness and Driving Continual Improvement: Evidence-Based Risk Reduction.](#)
5. [PDA PAC iAM 2017 Survey on Post-Approval Change: Is the Regulatory Environment Hindering Much-Needed Innovation in the Pharma Industry](#)
6. [PDA Technical Report 68: Risk-Based Approach for Prevention and Management of Drug Shortages](#)

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