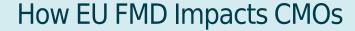




RESOURCES

Home Resources Resource Center





This two-minute video will give you a quick introduction to what the EU Falsified Medicines Directive (FMD) means for contract manufacturing organizations (CMOs), and what you need to do by the time it comes into force on the 9th of February 2019.

VideoEuropean Union Falsified Medicines DirectiveGlobal Track & TraceRegulatory/ComplianceEuropean Union

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A Guide to EU FMD for CMOs: Compliance, Collaboration and Business Opportunity

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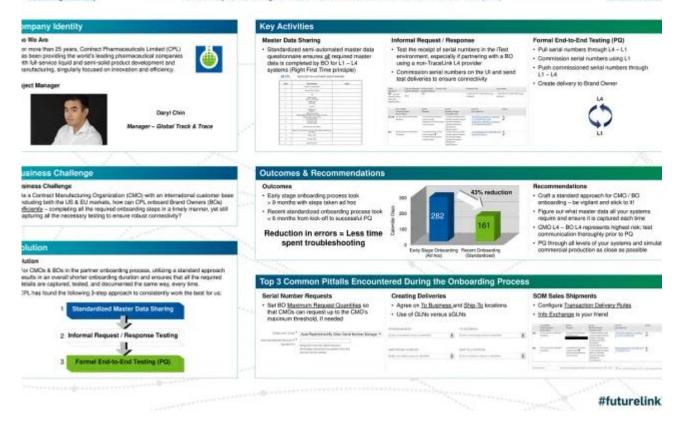


The CMO Serialization Perspective Utilizing a Standardized Approach for Efficient Partner Onboarding

CPL Outsourcing Made Easy

Author: Daryl Chin, Manager - Global Track & Trace, Contract Pharmaceuticals Limited (CPL)





Case Study: CPL | The CMO Serialization Perspective—Utilizing a Standardized Approach for Efficient Partner Onboarding

See how contract manufacturer Contract Pharmaceuticals Limited implemented a 3-step process for smooth pharmaceutical partner onboarding.





Can My CMO Upload Product Data to the European Hub?

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Stockpiling product to delay EU FMD compliance is a temporary approach that offers no long-term value—and carries significant risks.





Serialization: The Power to Transform Your Business

See how serialization helps you achieve compliance, prevent delays, protect your brand, and provide greater visibility into your products and patients.





Senior Vice President of Proc

Preparing for Saudi Arabia Compliance Reporting

TraceLink SVP of Product Marketing, Lucy Deus, provides an in-depth review of the January 2019 Saudi reporting requirements for pharma supply chains.