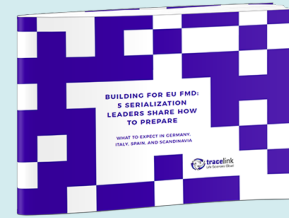




RESOURCES

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Building for EU FMD - 5 Serialization Leaders Share How to Prepare



As your pharmaceutical company builds and deploys serialization and track and trace capabilities to support the EU Falsified Medicines Directive (EU FMD), the challenge of managing all your requirements and specifications will become more and more complex. And time-consuming.

To help you prepare, we asked select EU serialization experts to share their views and lessons learned about overcoming implementation hurdles.

Learn:

- How frontrunners across Germany, Spain, Scandinavia, and Italy are addressing EU FMD.
- What the biggest roadblocks are right now.
- What makes implementing Level 1-Level 5 so complex.
- How you can improve compliance outcomes as the February 9, 2019, deadline gets closer.

If you have additional questions, we invite you to [marketing \[at\] tracelink.com](mailto:marketing@tracelink.com) (Subject: EU%20FMD%20Questions) (contact us) and speak with a serialization expert.

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More Serialization and Compliance Resources



An Interview with Tjoapack: Innovation through Serialization

Learn how Tjoapack turned the challenge of updating packaging for EU FMD into an opportunity for innovation, in this on-demand webinar.

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EU FMD, Distribution, and Decommissioning: Top Questions Answered

Learn the answers to seven decommissioning questions surrounding EU FMD and warehouses.

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How Santen Pharmaceutical is Using Serialization to Drive Supply Chain Transformation

Frank Binder, head of global supply chain management at Santen, discusses how serialization will allow Santen to gain operational intelligence and drive growth.

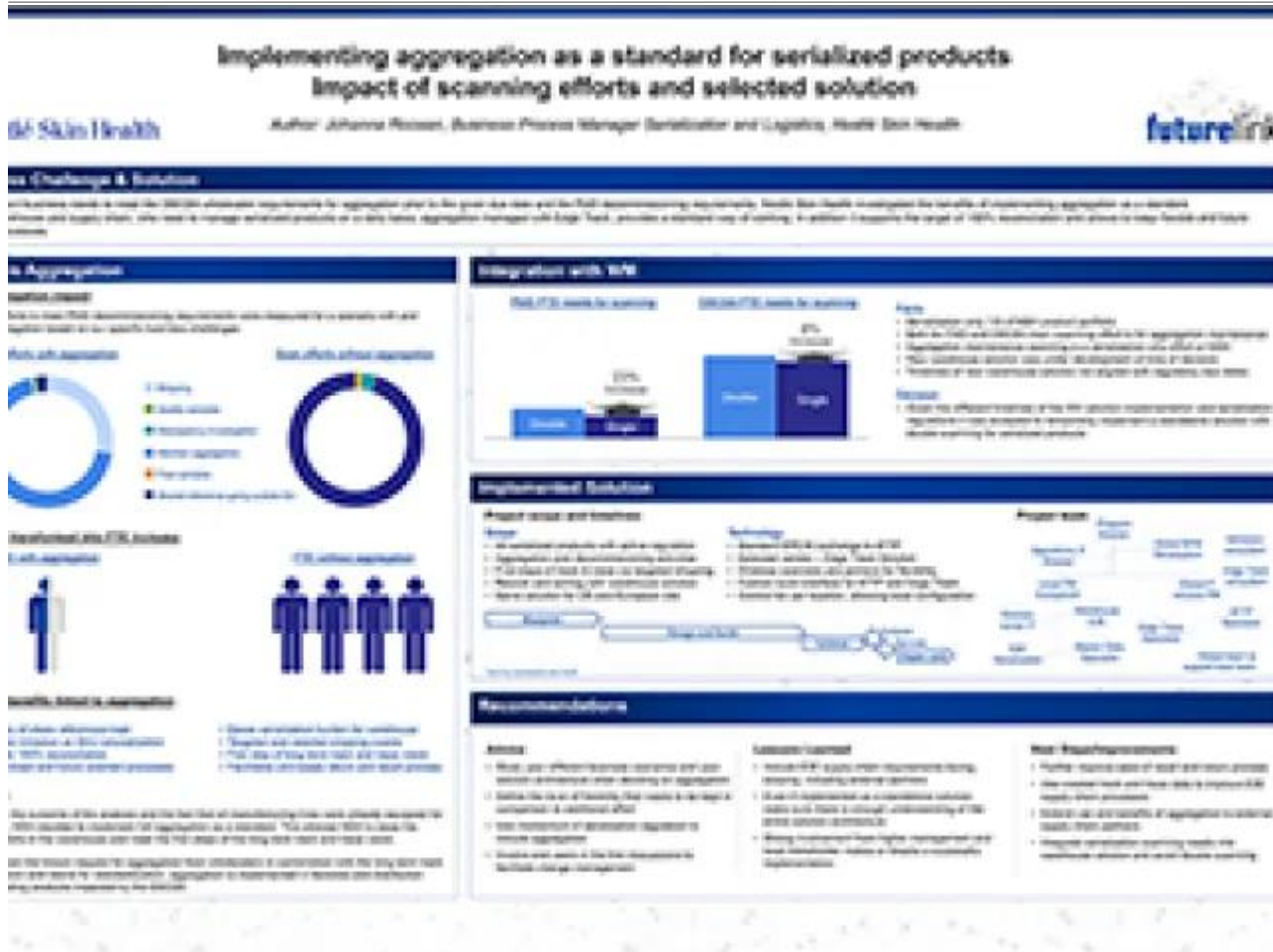
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Laetus Q&A: How People, Process, and Planning Will Drive Your EU FMD Compliance Success

Track and Trace Manager at Laetus explains why humans have a major impact on traceability, and what insights supply chain companies can gain from serialization.

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Learn how Nestlé Skin Health uses Edge Track to support their aggregation goals.

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ACS Q&A: Building a National System for EU FMD

Read an interview with ACS PharmaProtect to learn how the German national system works and how it will operate under EU FMD.

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