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Getting Started: EU FMD Guide to Pharma Serialization



If your company is part of the pharmaceutical supply chain in Europe, preparing for the EU Falsified Medicines Directive should now be a priority. The regulation presents considerable challenges and responsibilities—serialization, government reporting, verification—so understanding the requirements and knowing how to prepare for the February 2019 deadline is a must.

Our latest infographic can get you started on the road to EU FMD compliance. It covers the regulations' essential facts, including which nations are affected—and which have a compliance grace period—what each supply chain company must do to comply, and other vital details about:

- Which products can be grandfathered.

- The 2D DataMatrix code format.
- The specific data that must be uploaded and maintained.

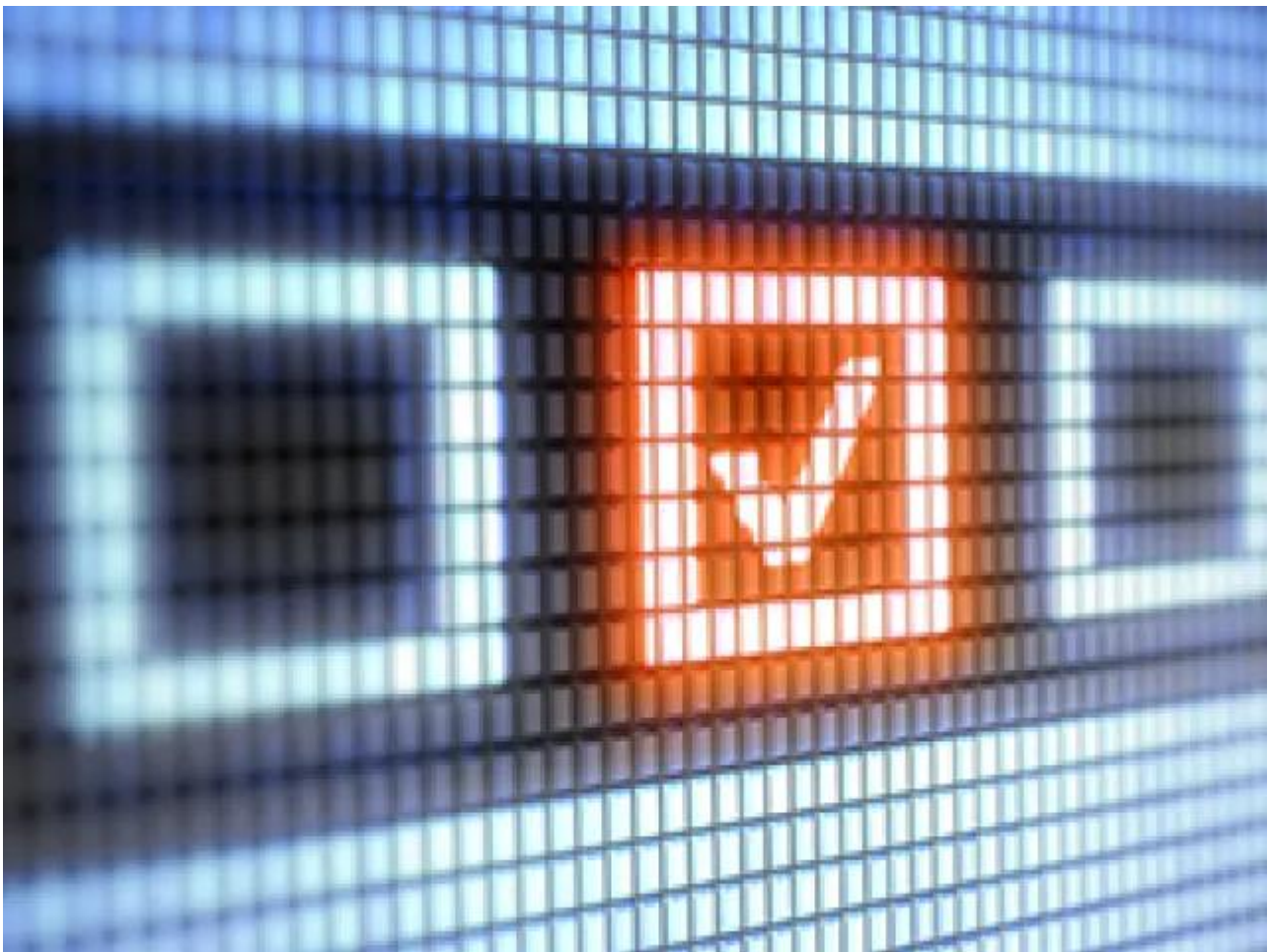
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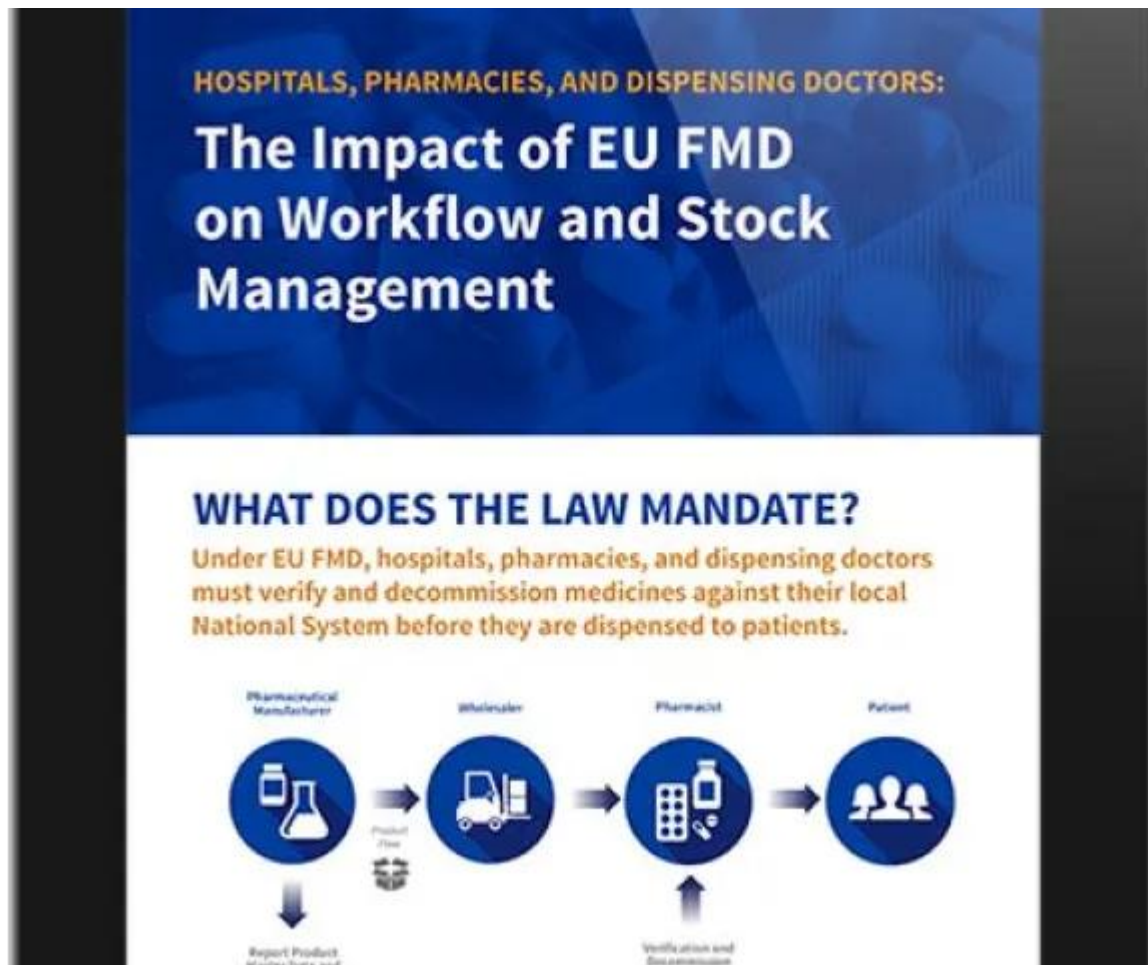
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Can You Afford a Manual Approach to EU FMD Compliance?

Manual data upload through the EMVO portal is simple in concept, but preparing and maintaining that data can be a challenge for smaller companies.

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The Impact of EU FMD on Pharmacy Workflow and Stock Management

View an infographic on how EU FMD changes hospital and pharmacy workflows.

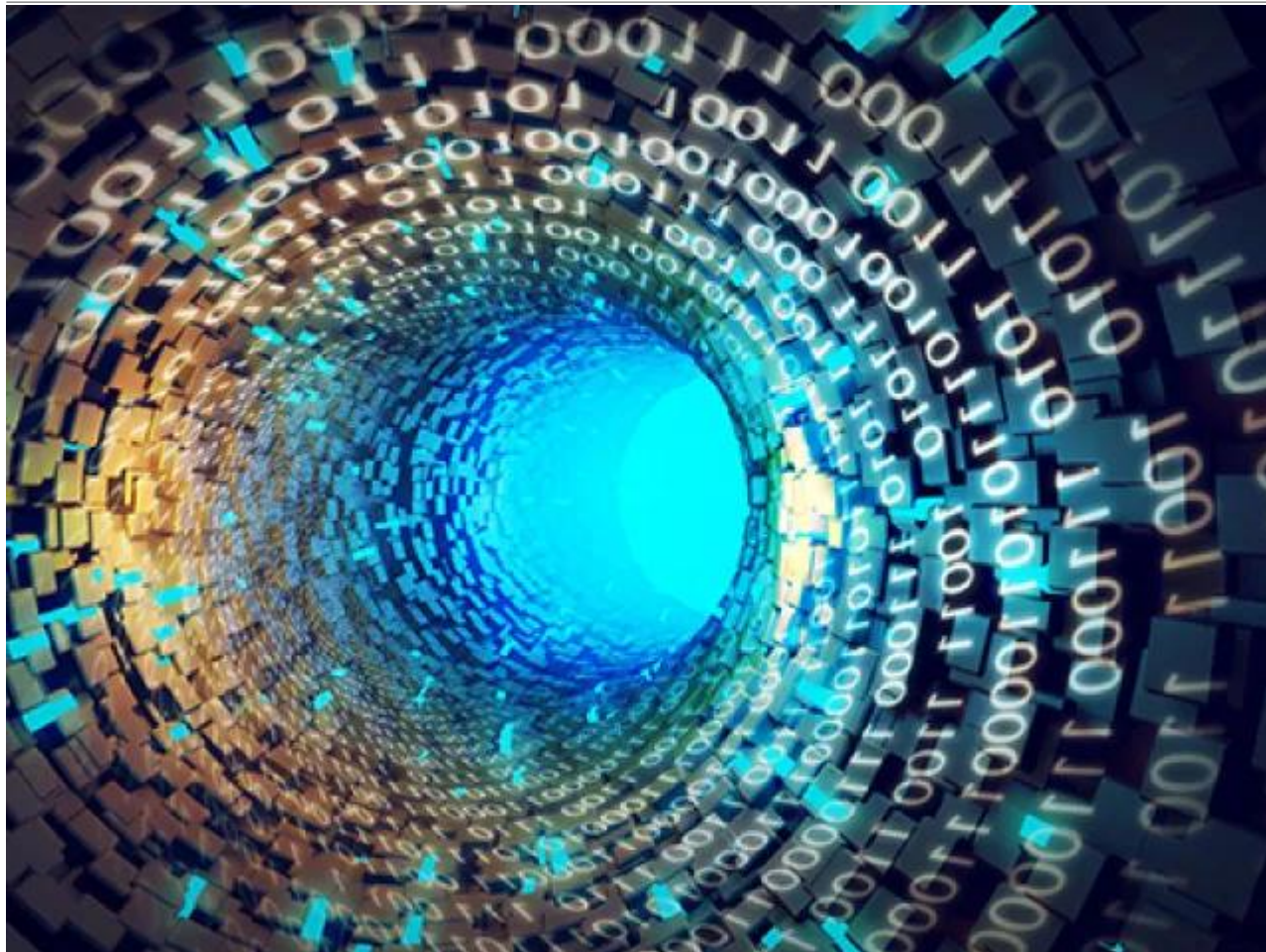
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What Are the 3 Major Requirements of EU FMD?

The EU Falsified Medicines Directive contains requirements for safety features and verification that details how companies must establish serialization and reporting.

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EU FMD Express: The Perfect Partner for Virtual Pharma

Discover why this small French pharma company chose TraceLink for a cost-effective compliance solution.

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Savior Lifetec: Serialization-Ready on TraceLink's Network

Find out how TraceLink partnered with Savior Lifetec to serialize its product line and ultimately increase patient safety.

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Can My CMO Upload Product Data to the European Hub?

Learn whether or not contract manufacturing organizations can upload product data to the European Hub for pharma companies.

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