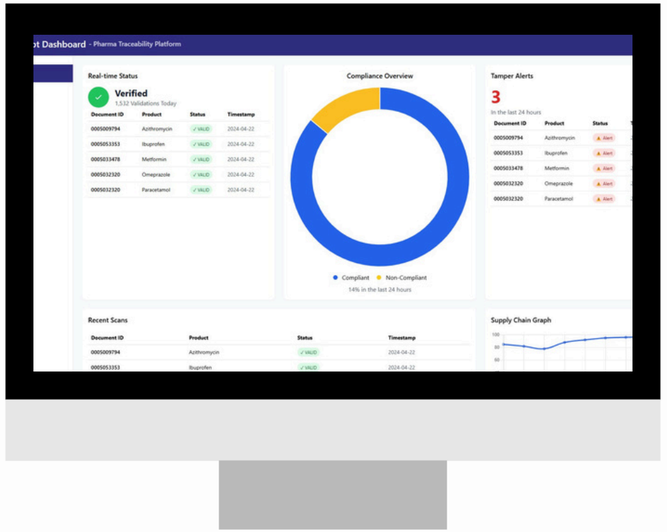
2026 Compliance Checklist for Pharma QA Managers

Ensure your packaging, traceability, and documentation are audit-ready—every day of the year.

Download Full Compliance Guide



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Is Your Pharma Chain Audit-Ready?

Intoday’s pharmaceutical landscape,QA leaders must navigate increasingly complex requirements—while maintaining the speed and safety of the supply chain.

From serialization and traceability to tamper-proof packaging and data readiness, the margin for error is shrinking. This practical checklist is designed to help you get ahead of compliance, protect your company’s reputation, and build a process your regulators will trust.

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Serialization Standards

- ☑ **Ensure GTIN, Expiry Date, Batch Number & Serial Number are human- and machine-readable**
- ☑ **Serialization formats must comply with DSCSA, EU FMD, or applicable region**
- ☑ **Unique product identifiers generated and verified before market release**
- ☑ **Serialized codes integrated with track-and-trace platform**

Tip: Use blockchain-linked serialization to protect against code cloning.

Batch Logging & Digital Traceability

- ✓ End-to-end tracking of every unit from production to pharmacy
- ✓ Digital logs for each batch, with automated time-stamped checkpoints
- ✓ Ability to trace recalls or errors back to the source in < 60 seconds
- ✓ Cloud-based ledger with access control & version history

Use real-time dashboards to flag missing data before audit day

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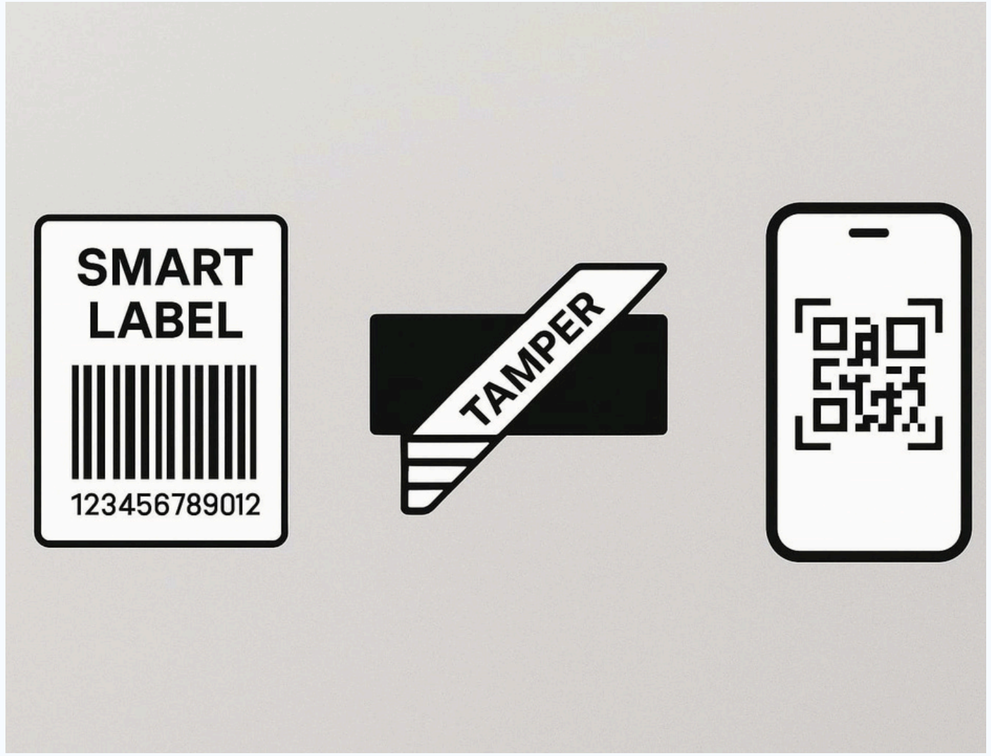
Tamper-Proof Packaging Protocols

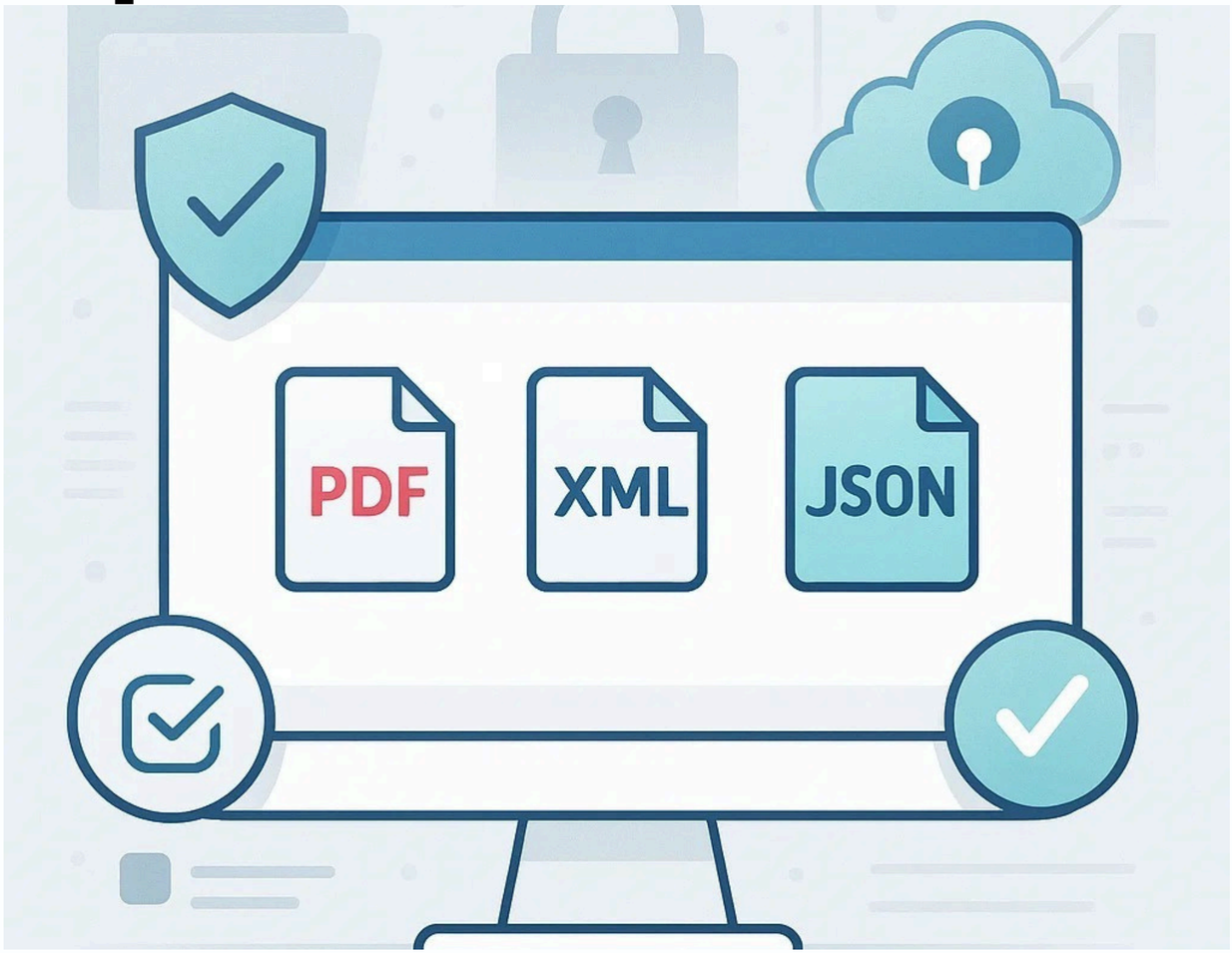


- ✓ Tamper-evident seals on every consumer-facing package
- ✓ Use of irreversible labels, tear indicators, or dual-seal layers
- ✓ Packaging verification through QR code scan or NFC tap
- ✓ Compliance with ISO 21976:2018 (Tamper Verification Features for Packaging)

Tip: Smart packaging adds traceability without increasing complexity

Data Storage, Regulatory Formats & Audit Preparedness

- ☑ Audit-ready reports exportable in XML, PDF, and JSON
- ☑ Role-based access to sensitive compliance logs
- ☑ Integration with FDA, EMA, CDSCO reporting portals
- ☑ Annual audit simulation run completed with QA team



Tip: Automate compliance documentation exports with scheduled reports.

Your Compliance Hub — Live, Logged & Verified

Synchrypt'sreal-time compliancetoolsgiveQA teams everything theyneedtomeetglobal pharma regulations—withoutchasing paperworkorriskingmissed errors.
Let your systems track what matters, while your team focuses on quality.

Ready to simplify compliance?
Book a free demo with our product specialist and explore how Synchrypt supports serialization, traceability, tamper detection, and reporting—in one unified system.
Visit: www.synchrypt.com

Book a Demo

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