Medcrypt

SBOM Regulatory Preview: Streamlined Compliance with Medcrypt's Helm

As SBOM (Software Bill of Materials) requirements become increasingly critical for medical device security and compliance, ensuring comprehensive documentation is key to efficient development cycles. Medcrypt's SBOM Regulatory Preview, powered by our Helm Vulnerability Management platform and regulatory experts, offers medical device manufacturers an expert review backed by our extensive industry experience. Our team of medical device security experts has helped manufacturers successfully navigate regulatory submissions. Without purchasing a full license, your team can now leverage our proven technical expertise and tooling to ensure your SBOM meets industry best practices and technical standards.

Regulatory Preview Program Overview

1. **Submit Your SBOM:** We accept CycloneDX, SPDX and Excel
2. **We process your SBOM:** We'll scan it for vulnerabilities and compliance gaps
3. **Regulatory style Report:** Receive a report mirroring an FDA hold letter, highlighting potential issues that could delay approval.

Proactive Issue Resolution

Identify and address potential issues before submission, minimizing costly resubmissions and delays.

Accelerate Submissions

Ensure your SBOM meets FDA requirements on the first pass, expediting the approval process and reducing time-to-market.

Revenue Protection

Hit projected launch dates and prevent the revenue loss associated with delays and prolonged regulatory review.

Secure Compliance, Accelerate Market Entry.

Reach out to get started on your SBOM Regulatory Preview and give your team the advantage of confidence and efficiency.

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