

UNLOCKING SUCCESS:

The Power of eQMS

In the complex MedTech landscape, Greenlight Guru's Electronic Quality Management System (eQMS) stands as a strategic solution. Our eQMS is tailored to drive efficiency, mitigate risk, and expedite innovation, positioning your organization as an industry leader. An independent study by Hobson & Company reveals that our eQMS solution delivers an ROI of 419% over the course of the contract, with a payback period of just 2.7 months. Partner with us to transform operational excellence.

The Crucial Need for eQMS

In today's highly regulated and rapidly advancing MedTech industry, relying on manual or paper-based quality management systems is no longer viable. The increased complexity in regulatory requirements, coupled with the urgency to accelerate innovation, requires a transition to an Electronic Quality Management System (eQMS). Failing to do so can lead to delays in time-to-market, regulatory compliance failures, inefficiencies in resource utilization, and potentially losing your competitive edge. With the stakes higher than ever, the adoption of an eQMS is not merely an option but a strategic necessity.

Strategic Alignment for MedTech Leaders

Greenlight Guru's eQMS is more than a technological advancement; it's a prudent decision for MedTech leaders aiming for efficiency, speed, and reduced risk. By embracing this solution, you're equipping your organization with the agility required in an innovation-driven market.

Key Benefits



reduction in time
needed to set up a QMS



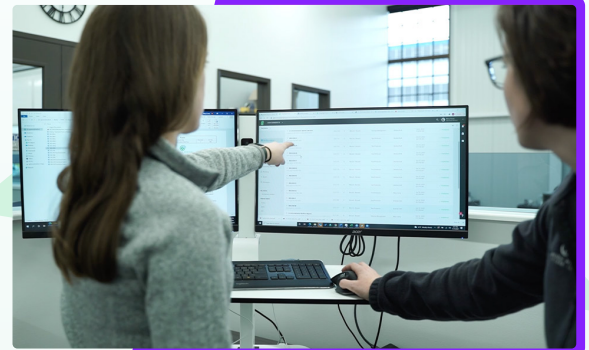
reduction in time
to market



reduction in time
spent on development/
design documentation

“ **What we pay to have Greenlight Guru is repaid in efficiency.”**

Clive Seymour,
CEO of Canterbury Scientific



Why Choose Greenlight Guru for Your eQMS

Built for MedTech

- ✓ **Built-in Compliance:** Tailored workflows for regulations like FDA 21 CFR Part 820, ISO 13485:2016, and EU MDR
- ✓ **Audit and inspection Readiness:** Industry-specific quality management software aligns with current medical device regulations, enhancing audit readiness and yielding unprecedented visibility
- ✓ **Design Control & Risk Management Integration:** Identify and mitigate risks early in development
- ✓ **Effortless Part 11 Validation:** Automated, risk-based approach simplifies validation, assuring compliance with evolving requirements

Gain Efficiency, Save Time

- ✓ **Efficiency and Effectiveness:** Saves substantial project hours during design, boosting efficiency up to 35%
- ✓ **Accelerated Implementation:** Support scaling without compromising compliance
- ✓ **Time-Saving Operational Efficiencies:** Enhances workflow and information management efficiency
- ✓ **Training Efficiency:** 75% reduction in QA/RA FTE time spent on mandatory training

Enable Growth

- ✓ **Enhanced Oversight and Consistency:** Leveraging a close-loop quality system, you gain real-time visibility into your QMS, ensuring informed decision-making, streamlined process management, and reduced non-compliance risks
- ✓ **Advance Skills with Greenlight Guru Academy:** Over 300 curated courses, certificates, and industry insights
- ✓ **User-friendly Visual QMS:** Easy-to-use platform that's ready out of the box
- ✓ **Best-in-Class EDC Solutions:** Facilitates high-quality clinical trials

**LEADER****QMS SOFTWARE**

Since Winter 2019

MEDICAL QMS SOFTWARE

Since Spring 2021



Your Strategic Move

Embracing Greenlight Guru as your eQMS is not just a technological decision: it's a strategic business move. Ensure your organization stays ahead of the curve, compliant, and primed for innovation. Bring your medical device to market safer and faster.