

Ensure product quality and patient safety with the **right Quality Management System (QMS)**

Save Time and Boost Your Bottom Line

Ignoring Your QMS Could Cost You

Discounting the development and implementation of an effective QMS can be a costly mistake. This short-sightedness can result in unnecessary costs, delays to market, and reallocation of valuable internal resources to manage, investigate, and correct errors that could be avoided from the start.

Eliminate the wasted time and expense spent on workarounds and corrections. Partner with the industry-leading compliance experts to evaluate your QMS and identify issues before incidents occur. Fewer incidents translates to added capacity, increased productivity, faster profit returns, and the opportunity for your valued resources to focus on additional product development.

Efficient and Effective QMS:

- Ensures you are operating in a compliant manner
- Bridges your organization's compliance practices with regulatory requirements
- Establishes the infrastructure and oversight requirements for manufacturing drugs, medical devices, and APIs under Good Manufacturing Practice (GMP), for conducting clinical trials in the form of Good Clinical Practice (GCP), and for performing clinical laboratory testing as Good Clinical Laboratory Practices (GCLP)
- Creates a platform for maintaining consistency across dynamic regulated environments throughout the product lifecycle



Experienced Partners for Development and Implementation

ProPharma Group is THE trusted partner in designing, building, and managing risk-based and phase-appropriate QMS programs. Our team of industry-leading experts understands the complex regulatory environment and the various compliance components that are required for your organization to succeed. We closely collaborate with your team and help optimize the processes specific to your operational requirements and quality programs.

At a minimum, our aim is to ensure your QMS achieves three objectives:

- Supports continuous improvement
- Meets all regulatory requirements and expectations
- Establishes risk-based and “right-sized” phase appropriate processes

Looking to develop and implement a new QMS? Need to optimize your current QMS? Our team of compliance and quality experts can help you develop, implement and manage the right QMS that ensures product quality and patient safety.

Improving Patient Health and Safety. **At Every Step.**

From early concept development through each clinical phase, product launch, and commercialization, we partner with pharmaceutical, biotechnology, and medical device clients to tackle complex challenges. We help to ensure regulatory expectations are met, business goals are achieved, and patient health and safety is improved.

Contact ProPharma Group today for a complete evaluation and explore your options to perfect your QMS.

 www.ProPharmaGroup.com

 Info@ProPharmaGroup.com



Improving Patient Health and Safety

