Enzyme QMS

Overview



Whether you're in the market for your first QMS, looking to move away from a paper-based system, or have outgrown your current QMS, Enzyme has everything you need, premarket to postmarket. Designed to grow and evolve as you do, Enzyme QMS empowers you to keep pace with innovation while maintaining compliance globally.

- ☑ 21 CFR Part 11 compliant
- ☑ Validated out-of-box
- ☑ Built for medical device, digital health, and biopharma companies
- ☑ Customizable to you
- ☑ Offers an industry-leading number of integrations

Plans

Core

- **Document control**
- **Training**
- **eSignatures**
- Design control
- Risk management
- Suppliers
- Audits
- Complaints
- Nonconformance
- CAPA

Premarket

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Postmarket

- **Document control**
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- **eSignatures**
- **Design control**
- **Risk management**
- **Suppliers**
- **Audits**
- **Complaints**
- **Nonconformance**
- **CAPA**

In-house consulting services are also available to help with everything from day-to-day support, to QMS optimization, to writing submissions and acting as your representative to notified bodies.

Pricing

Our pricing packages are based on the plan you choose and the number and types of users you need. We offer two types of users: Full and Lite. Lite is for users who only need to view and sign off on training. Discounts are also available based on length of term, payment frequency, and contract start date.

Enzyme

Serving teams from 2 to 2,000, premarket to postmarket, startup to enterprise, Enzyme is the only QMS you'll ever need — or want.