

Stay ahead. Stay compliant. With **AIMEDIQ**  
The AI platform for regulatory clarity and control in MedTech.

**The challenge****Regulatory overload is real.**

MedTech companies face a constant flood of changing laws, standards and guidelines across multiple markets. Manual tracking is inefficient, time-consuming, and prone to error – and missing a single update can have serious consequences.

**The key****AIMEDIQ makes compliance easy.**

Swiss-engineered and AI-powered by DocDive from G+D Netcetera, AIMEDIQ simplifies and accelerates access to essential regulatory information – all in one place. It's built for MedTech professionals who want clarity, structure and peace of mind.

**What it delivers:**

- One centralised, searchable platform
- 1,600+ official documents: laws, directives, and guidelines
- Global coverage (EU, UK, US, CH, Japan, Singapore, Australia – more to come)
- AI-curated responses – fast, tailored, complete

**Key features:**

**Real-time updates:** regulatory changes and user-specific standards

**Global coverage:** EU, UK, US, CH, Japan, Singapore, Australia

**Smart alerts:** timely expiry notifications (planned)

**Unlimited access:** flat-rate licence with no query limits

**Custom uploads:** manage own licensed standards and files

**Tailored results:** AI adapts to your standards and content

**Document traceability:** full transparency of all sources

**Cloud-based:** secure, closed AWS system in Frankfurt

**What makes AIMEDIQ different****Stay ahead of regulatory changes**

Continuous updates from trusted regulatory sources

**Effortless compliance**

Reduce errors with smart automation

**Save time**

Focus on high-value tasks, not standard maintenance

**Scalable & secure**

Works for start-ups and global MedTech leaders alike

**Complete visibility**

All relevant documents connected, even with vague search terms

**Expiry monitoring**

Get notified before documents expire

Live demos available!

**Ready to simplify compliance?  
Reach out to Mike Seidenberg:**

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