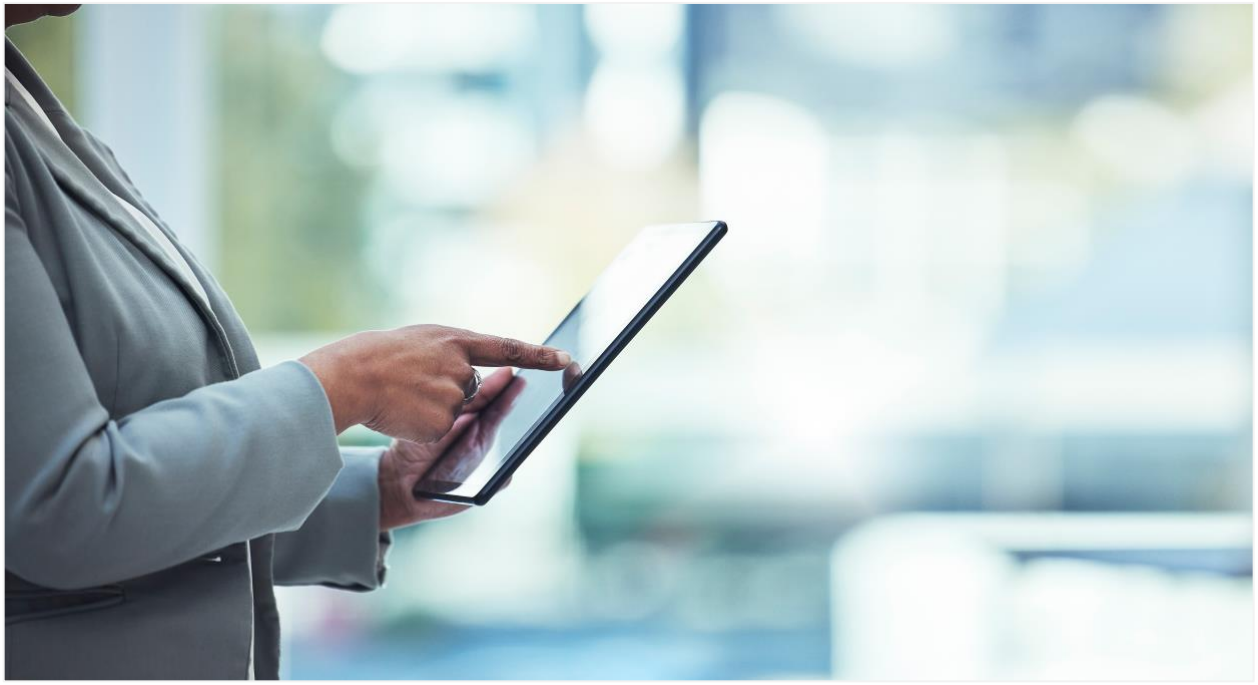
Revolutionize Your Quality Management and Regulatory Compliance



Welcome to **Ryden**, your all-in-one solution for transforming how MedTech companies manage and maintain quality and regulatory compliance. Our state-of-the-art **SaaS platform** harnesses the power of **AI** to streamline processes, reduce complexity, and deliver actionable insights across your organization.

Call us at 1.312.219.5471 or email at sales@ryden.ai

Why Ryden?

AI-Driven Features:
We leverage advanced AI trained for real-time regulatory gap analysis. Whether you're a startup or a global enterprise, Ryden helps you stay proactive and informed.

Scalable, Multi-Facility Architecture: Built with large organizations in mind, our platform ensures each user experiences a custom-tailored environment—perfect for multi-site and multi-country operations.

Customizable for Global Compliance: With Ryden, you can choose laws, regulations, and industry standards to meet the needs of your compliance requirements. Our intelligent recommendations adapt to evolving global regulatory landscapes.

Key Objectives & Goals

1. **AI-Powered Compliance Assessments**
   - Simplify complex regulatory requirements.
   - Receive automated gap analyses with clear, actionable insights.
2. **Holistic Quality Maturity Evaluation**
   - Identify strengths and weaknesses at both company and facility levels.
   - Benchmark performance over time against industry standards.
3. **Project Management for Compliance**
   - Assign tasks, track progress, and compile final compliance reports.
   - Collaborate across departments and regions to maintain alignment.

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How It Works

- **Law Assessment**: Input your target laws, and Ryden's AI will assess, categorize, and flag requirements relevant to your product lines.
- **Device/Pharma Project Requirements**: Address unique compliance needs for medical devices or pharmaceuticals—Ryden tracks all pertinent regulations in one place.
- **Processes and Procedures**: Build out your organizational best practices with SOPs and policies that align with global standards.

Gap Analysis Made Easy

Ryden works with your essential documentation to keep teams synchronized and compliant:

1. **Manuals/Policies (1–5)**
    - High-level documents defining device and legal compliance.
    - Essential for establishing a strong regulatory framework.
2. **Standard Operating Procedures (SOPs) (5–100)**
    - High-level processes ensuring quality and safety.
    - Aimed at streamlining daily operations across multiple facilities.
3. **Work Instructions (0–100)**
    - Step-by-step guidelines tailored for specific tasks.
    - Ideal for training staff and reducing errors in production.
4. **Form Templates (0–100)**
    - Standardized data capture for audits, inspections, and more.
    - Helps teams stay organized while minimizing administrative burden.

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QMS Document Integration

Ryden works with your preferred QMS system, and will seamlessly integrate and automate the processing of documents:

- **Policies/Manuals**: Define your device's regulatory scope and legal obligations.
- **Procedures**: Implement high-level processes to meet compliance objectives.
- **Instructions/Form Templates**: Provide the granular detail your teams need to execute tasks accurately and consistently.

Your Path to Seamless Compliance

Ryden works with your essential documentation to keep teams synchronized and compliant:

1. **Analyze** – Feed your regulatory requirements into Ryden's AI engine for immediate gap analysis and best practice assessments.
2. **Implement** – Assign tasks through our project management tools for SOPs, forms, or instructions tailored to specific compliance goals.
3. **Monitor & Improve** – Track real-time metrics on your organization's compliance health and compare performance against industry standards to continuously drive improvement.

Start Your Journey with Ryden

By choosing Ryden, you're not just investing in a platform—you're empowering your organization to excel in **quality management** and **regulatory compliance**. Whether you're targeting a single market or scaling globally, Ryden's flexible, multi-lingual, AI-driven solution is designed to meet your needs today and adapt to your challenges tomorrow.

Ready to Transform Your Compliance Process?

Contact us at sales@ryden.ai or visit ryden.ai to learn how Ryden can revolutionize the way you manage quality and regulatory compliance. Let's build a future of simpler, smarter, and more secure compliance—together.