

If You Don't Understand It, You Can't Translate It!

At a glance

Quality Management System (QMS) documentation defines processes, policies, and responsibilities essential for ensuring medical device quality, safety, and regulatory compliance globally. Accurate translations streamline international compliance, optimize operational efficiency, and enhance audit readiness.

Key metrics



35%

Reduction in global audit findings and compliance discrepancies



99.7%

On Time Delivery



100%

Regulatory Compliance

CHALLENGES



QMS documents require precise translation of detailed regulatory guidelines, procedural instructions, and compliance standards. Translation errors or inconsistencies can lead to regulatory audits, compliance issues, procedural deviations, or operational inefficiencies. Translating extensive, technical QMS content consistently across multiple languages and regulatory environments adds significant complexity, especially within tight operational and regulatory timelines.

SOLUTIONS



AI-optimized translations combined with expert medical device regulatory linguists ensure precise, compliant translations of QMS documentation worldwide.



**Quality and Compliance
Specialists**



**12 Step Quality
Control Process**



**Scalable To Meet
Your Needs**

BENEFITS



1

Enhanced Global Compliance

Accurate translations standardize quality processes internationally, significantly reducing audit findings and compliance risks.

2

Improved Operational Consistency

Clear, consistent QMS translations simplify global operations, optimize training, and reduce procedural deviations.

3

Reduced Regulatory Risk

Reliable translations prevent costly compliance issues, audits, and disruptions, safeguarding global market operations and reputation.