

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

## At a glance

Risk assessment reports systematically identify, evaluate, and document potential hazards associated with medical devices. Accurate translations ensure clear global communication of device-related risks, support regulatory compliance, enhance patient safety, and streamline international market approvals.

### Key metrics



**40%**

Reduction in regulatory approval timelines



**99.7%**

On Time Delivery



**100%**

Regulatory Compliance

## CHALLENGES



Risk assessment reports include complex technical details and precise regulatory language requiring exact translations. Translation inaccuracies or ambiguities can compromise patient safety, trigger regulatory scrutiny, delay product approvals, or result in costly market recalls. Translating detailed risk assessments clearly and consistently across multiple languages and regulatory jurisdictions adds significant complexity, especially under tight submission deadlines.

## SOLUTIONS



AI-optimized translations combined with specialized medical device risk linguists ensure accurate, compliant translations of risk assessment reports globally.



**Risk Assessment Specialists**



**12 Step Quality Control Process**



**Scalable To Meet Your Needs**

## BENEFITS



**1**

### Enhanced Patient Safety

Accurate translations ensure clear communication of potential device risks, significantly reducing patient safety incidents globally.

**2**

### Improved Regulatory Compliance

Compliant translations streamline regulatory acceptance, minimizing audits, queries, and approval delays.

**3**

### Reduced Market Entry Risks

Precise translations protect against costly regulatory actions and market disruptions, safeguarding global commercialization strategies.