

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

## At a glance

Clinical trial sponsors and research organizations rely on precise translations of site-specific operating procedures, guidelines, and manuals found in site Master Files and clinical site documentation. Accurate translations ensure regulatory compliance, standardize global site operations, and support trial consistency and efficiency.

## Key metrics



**>30%**

Reduction in Clinical Site Compliance Discrepancies



**99.7%**

On Time Delivery



**100%**

Regulatory Compliance

## CHALLENGES



Site operations documents contain detailed clinical and regulatory instructions requiring precise translations. Any inaccuracies can disrupt global clinical consistency, regulatory compliance, and site operations. Ensuring clarity, uniformity, and adherence to strict regulatory standards across multiple languages adds complexity, especially under tight deadlines.

## SOLUTIONS



AI-optimized translations combined with expert clinical and regulatory linguists ensure site operations documents maintain accuracy, compliance, and operational consistency globally.



**Clinical Expertise**



**12 Step Quality Control Process**



**Scalable To Meet Your Needs**

## BENEFITS



**1**

### Standardized Global Operations

Accurate translations ensure operational consistency, improving site adherence to protocols and reducing deviations globally.

**2**

### Enhanced Regulatory Compliance

Precise translations minimize regulatory queries, audits, and operational discrepancies, improving site inspection readiness.

**3**

### Streamlined Site Management

Rapid, accurate translations simplify site training, onboarding, and operations, increasing overall clinical trial efficiency.