

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

At a glance

Case Report Forms (CRFs) collect essential clinical trial data on medical device performance, patient outcomes, and safety events. Accurate translations ensure reliable data collection, regulatory compliance, improved clinical outcomes, and efficient global trial management.

Key metrics



40%

Reduction in clinical data queries



99.7%

On Time Delivery



100%

Regulatory Compliance

CHALLENGES



CRFs contain detailed clinical, procedural, and safety information requiring precise translations. Errors or unclear wording can result in inaccurate data collection, regulatory submission delays, compromised patient safety monitoring, and invalid trial outcomes. Translating CRFs consistently and clearly across multiple languages and regulatory jurisdictions, particularly within tight clinical timelines, adds significant complexity to the global clinical trials.

SOLUTIONS



AI-optimized translations combined with specialized clinical linguists ensure accurate, compliant translations of CRFs globally.



**Clinical Data
Experts**



**12 Step Quality
Control Process**



**Scalable To Meet
Your Needs**

BENEFITS



1

Enhanced Clinical Data Accuracy

Precise translations significantly improve data reliability, supporting clear interpretation of clinical outcomes globally.

2

Improved Patient Safety Monitoring

Accurate CRFs enable consistent and reliable safety reporting, enhancing global patient protection efforts.

3

Accelerated Regulatory Approvals

Compliant translations streamline regulatory reviews, facilitating quicker approvals and timely global market entry.



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