



Patient Source Documents, Admission and Discharge, Labs Use Case

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

At a glance

Accurate translation of patient source documents—including admission records, discharge summaries, and laboratory reports—is vital for clinical data integrity, patient safety, and regulatory compliance. Precise translations ensure seamless data management, support global clinical trials, and facilitate timely regulatory submissions.

Key metrics



>40%

Reduction in Data Queries and Corrections



99.7%

On Time Delivery



100%

Regulatory Compliance

CHALLENGES



Patient source documents such as admission records, discharge summaries, and laboratory reports contain sensitive, complex medical data requiring precise translations. Errors can result in misinterpretations, compromised patient safety, data discrepancies, or regulatory issues. Translating highly varied clinical documents consistently and accurately under strict clinical timelines adds further complexity to global clinical operations.

SOLUTIONS



AI-optimized translations complemented by specialized clinical linguists ensure patient source documents accurately reflect the original clinical data, maintaining regulatory compliance and data integrity.



Clinical Trial Expertise



12 Step Quality Control Process



Scalable To Meet Your Needs

BENEFITS



1

Improved Clinical Data Accuracy

Accurate translations ensure consistent patient data across multiple regions, reducing clinical discrepancies.

2

Regulatory Confidence and Compliance

High-quality translations minimize regulatory queries and risks, streamlining submissions and audits.

3

Enhanced Patient Safety and Care

Clear, error-free translations protect patient safety and facilitate better clinical decision-making globally.



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