

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

At a glance

Informed Consent Forms (ICFs) are essential documents in clinical trials, clearly communicating risks, benefits, and procedures to participants. Accurate translation ensures patient understanding, compliance with ethical standards, and successful regulatory approvals for multinational studies.

Key metrics



35%

Faster Patient Enrollment



99.7%

On Time Delivery



100%

Regulatory Compliance

CHALLENGES



Informed consent forms (ICFs) require precise translation that accurately conveys complex medical concepts, risks, and procedures in clear, understandable language for diverse patient populations. Misinterpretations can compromise patient safety, create ethical issues, and lead to regulatory delays. Maintaining cultural appropriateness and regulatory compliance across multiple languages adds complexity, particularly when meeting tight timelines for clinical trial initiation.

SOLUTIONS



AI-optimized translation processes combined with expert medical linguists ensure informed consent forms clearly communicate essential clinical information, ethical requirements, and patient rights accurately across languages.



Medical Experts



12 Step Quality Control Process



Scalable To Meet Your Needs

BENEFITS



1

Enhanced Patient Safety

Clear, precise translations enable patients to make informed decisions, ensuring ethical compliance and patient protection globally.

2

Streamlined Ethical Approval

Accurate translations reduce ethical committee queries and facilitate smoother regulatory approvals.

3

Improved Trial Recruitment

Patient-friendly translations increase participant comprehension and engagement, supporting faster enrollment and trial efficiency.