

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

At a glance

Pharmaceutical companies require precise translations of drug registration dossiers and data sheets to gain timely approval in global markets. Accurate, compliant translations streamline regulatory processes, enhance document consistency, and accelerate international product launches.

Key metrics



50%
Faster Regulatory Approvals



99.7%
On Time Delivery



100%
Regulatory Compliance

CHALLENGES

Drug registration data sheets include highly detailed scientific, clinical, and regulatory content requiring exact translations for successful regulatory submissions. Errors or inconsistencies can cause costly approval delays, regulatory queries, or market rejections. Meeting tight deadlines while maintaining accuracy and alignment with various international regulatory requirements further complicates this critical process.

SOLUTIONS

AI-optimized translations combined with experienced medical and scientific linguists ensure data sheets meet regulatory precision and consistency across languages and markets.



Regulatory Expertise



12 Step Quality Control Process



Scalable To Meet Your Needs

BENEFITS

1

Accelerated Regulatory Approvals

Accelerates drug registration through precise, compliant documentation, reducing regulatory hurdles.

2

Improved Global Compliance

Minimizes translation-related regulatory rejections, protecting market timelines and reputation.

3

Global Consistency

Ensures uniformity and precision across multilingual submissions, simplifying global dossier management.