

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

At a glance

Pharmaceutical companies depend on accurate translations of product dossiers and regulatory submissions to gain approval across global markets. Precise, compliant translations of dossier data sheets ensure successful submissions, enabling rapid market entry and consistent communication with international regulatory bodies.

Key metrics



50%
Faster Regulatory Approvals



99.7%
On Time Delivery



100%
Regulatory Compliance

CHALLENGES

Pharmaceutical data sheets within dossiers include extensive scientific, clinical, and regulatory content requiring flawless translation to satisfy global standards. Misinterpretations or inaccuracies can result in regulatory rejections, costly submission delays, or market entry setbacks. Ensuring translation consistency across multiple languages, while adhering to complex regional requirements and tight regulatory timelines, presents ongoing operational challenges.

SOLUTIONS

AI-optimized translations combined with expert linguist validation ensure dossiers are accurately translated to meet regulatory compliance across all target markets. Translation teams specialized in regulatory and scientific content accurately manage complex documents, maintaining consistency and precision.



Regulatory Expertise



12 Step Quality Control Process



Scalable To Meet Your Needs

BENEFITS

1

Accelerated Regulatory Approvals

Accurate translations facilitate faster dossier reviews, reducing approval timelines and enabling quicker global market entry.

2

Improved Global Compliance

Ensures dossiers align precisely with international standards, minimizing regulatory queries and rejections.

3

Enhanced Market Reach

High-quality translations expand pharmaceutical companies' ability to enter new markets quickly and confidently.