

Medical Device Product Labels and Packaging

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

At a glance

Accurate translation of medical device product labels and packaging is essential for patient safety, regulatory compliance, and successful international market entry.

Precise, compliant translations ensure clear communication of critical safety, usage, and regulatory information globally.

Key metrics



35%

Reduction in labelingrelated regulatory queries



99.7%
On Time Delivery



100%

Regulatory Compliance





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CHALLENGES



Medical device labels and packaging require precise translations of critical regulatory information, safety warnings, usage instructions, and symbols. Errors or inconsistencies can lead to misuse, patient harm, regulatory recalls, and market entry delays. Translating highly regulated content consistently and clearly across multiple languages and regulatory jurisdictions, often within tight commercialization timelines, adds complexity to the translation process

SOLUTIONS



Al-optimized translations combined with expert medical device regulatory linguists ensure accurate, compliant translations for labels and packaging.



Labeling Specialists



12 Step Quality Control Process



Scalable To Meet Your Needs

BENEFITS





Enhanced Patient Safety

Accurate, clear labeling translations minimize user errors and reduce the risk of patient harm.



Accelerated Regulatory Approval

Compliant translations streamline regulatory acceptance, expediting global product approvals and market access.



Reduced Compliance Risks

Reliable, consistent translations prevent regulatory actions, recalls, and costly market disruptions, protecting brand reputation.