



Medical Device Adverse Event Report

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

At a glance

Accurate translation of medical device adverse event reports is critical for timely global regulatory submissions, patient safety monitoring, and compliance with international vigilance requirements. Precise translations facilitate swift regulatory communication, prompt corrective actions, and effective risk management worldwide.

Key metrics



35%

Reduction in regulatory reporting delays



99.7%

On Time Delivery



100%

Regulatory Compliance

CHALLENGES



Adverse event reports contain urgent, complex clinical information and require immediate and exact translations. Errors can significantly impact patient safety, lead to regulatory noncompliance, market disruptions, or product recalls. Translating detailed adverse event information quickly and consistently across multiple languages, while strictly adhering to global regulatory requirements, adds complexity to the translation process

SOLUTIONS



AI-optimized translations combined with specialized medical device safety linguists ensure rapid, accurate, and compliant translations of adverse event reports.



**Safety Translation
Specialists**



**12 Step Quality
Control Process**



**Scalable To Meet
Your Needs**

BENEFITS



1

Accelerated Regulatory Reporting

Accurate, timely translations streamline adverse event submissions, ensuring compliance and preventing regulatory delays.

2

Enhanced Patient Safety

Precise translations clearly communicate device-related risks, facilitating swift and effective patient safety interventions globally.

3

Reduced Compliance Risks

Reliable translations mitigate costly regulatory actions, minimize audits, and protect market presence.



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