



Medical Device Post-Market Surveillance (PMS) Report

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

At a glance

Post-market surveillance (PMS) reports document ongoing device performance, safety, and risk management after medical device commercialization. Accurate translations ensure global regulatory compliance, consistent safety monitoring, and efficient communication of device performance internationally.

Key metrics



45%

Reduction in regulatory reporting delay



99.7%

On Time Delivery



100%

Regulatory Compliance

CHALLENGES



Post-market surveillance reports contain critical safety and clinical performance data requiring precise translations for global regulatory submissions. Inaccurate or unclear translations can compromise patient safety, trigger regulatory audits, and delay market approvals or renewals. Translating detailed, technical, and time-sensitive PMS documentation consistently across multiple languages adds complexity, particularly under strict reporting deadlines.

SOLUTIONS



AI-optimized translations combined with specialized medical device safety linguists ensure accurate, compliant translations of PMS reports worldwide.



Safety Translation Experts



12 Step Quality Control Process



Scalable To Meet Your Needs

BENEFITS



1

Enhanced Patient Safety Monitoring

Precise translations clearly communicate device risks and safety data, significantly improving global patient safety.

2

Improved Regulatory Confidence

Accurate, compliant PMS translations streamline regulatory reviews, minimizing audits and compliance issues.

3

Reduced Operational Risk

Reliable translations prevent costly regulatory queries, delays, and corrective actions, protecting market continuity and brand integrity.



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