Can better, faster, more accurate content really save your company $1 million a day?

Yes! According to a Univ. of Texas study, delayed drug approvals cost pharmaceutical companies $1 million a day. Understaffed or over-matched content development teams can contribute to these delays.

RWS offers a range of content development solutions to help customers deliver concise, multilingual content - at the highest standards of safety and quality - faster than ever.

Turnkey patient/clinician documentation support services

RWS content solutions help create technical documentation throughout the product lifecycle. This flexible offering ranges from source content optimization and occasional staff augmentation to turnkey outsourced documentation development and content strategy consulting.

- Source file creation & analysis including technical writing & illustration
- Content strategy, style and terminology guides, writer training
- Patient/clinician guides, IFUs, lay summaries, etc.
- Certificates of translation
- IFU analysis & MDR/IVDR compliance checks

We don't just translate IFUs into all needed target languages; we optimized the entire IFU creation process. After performing an IFU audit, our analysis allows us to:

- Verify documents for product liability and safety risks
- Increase brand adherence
- Lower translation and printing budgets
- Improve user experience

Our Turnkey Technical Writing and Translation Helps You Eliminate Risk

We know that creating high-quality, compliant technical documentation is a complex process. We now offer turnkey ISO-certified technical writing and translation capabilities for your technical documentation. We have designed specific services that assist our clients with successful MDR and IVDR compliance, audit readiness, and reduced translation budgets.

A Customer Success Story

We worked with a leading medical device company to update its IFU repository and translate the documents into 20+ languages in preparation for MDR and IVDR compliance.

As part of our process, we:

- Verified MDR & IVDR Compliance
- Handled the In-Country Review Process for the Client
- Optimized the IFU Creation and Translation Process
- Created and Verified Terminology

As a result, our client realized cost-saving, efficiency, and quality benefits, including:

50% Reduction In-Country Review Cycle Time

15% Reduction in Translation Budget

Created a **Consistent** and **Professional** IFU aesthetic for enhanced global branding and improved user experience

Increase MDR Compliance and audit readiness by identifying missing IFU content

Up to **25%** Reduction in Translation & Printing Budgets by Shortening IFUs

Consistently Translated Target Languages through terminology harmonization

We Work With

20 of the **Top 20**

Medical and In Vitro Diagnostic Device Companies

We Are

ISO Certified

ISO 9001:2015, ISO 13485:2016, ISO 17100:2015, ISO 27001:2013, ISO 18587:2016

We Have

30+ Years Experience

with exclusive focus on life sciences

For further information, please visit: rws.com/medical-devices