Solutions Overview

3 Ways to Safely Accelerate Your Clinical Document Submissions

PRIVACY ANALYTICS
an IQVIA company

Enabled by powerful privacy technology

What you're up against

As a clinical trial sponsor, you're facing pressure from multiple sides when it comes to handling regulatory requirements for document anonymization and redaction.

Meet tight timelines, ensure a competitive edge

To help you ensure a competitive edge, Privacy Analytics offers three paths to protecting CCI and personal data while meeting deadlines for EU CTR and other initiatives such as Health Canada PRCI and EMA Policy 0070: **Redaction-as-a-Service, Anonymization-as-a-Service, and licensed software**. Each solution is powered by our 4th-gen document redaction and anonymization software platform.

With Privacy Analytics by your side, you'll be better able to safely accelerate submissions for the broader range of clinical document types you now need to address throughout the clinical trial life cycle.

Reduce time to submit clinical trial documents from weeks to days

Our latest software release unites our proven natural language processing technology with automated redaction, providing you with a seamless solution to meet all of your evolving data privacy goals.

Which option is right for you?

License our software for your own use in-house, or rely on our team of 100+ privacy experts to leverage our software for you (freeing your team for other value-adding initiatives). Either way, you will accurately detect and protect sensitive data at scale, reducing the burden on your team.

	Redaction-as-a-Service	Anonymization-as-a-Service	Software licensing
What?	**Fast and cost-effective** way to handle time-sensitive requests.	**Industry best practice** approach for maximum transparency.	Use our software to efficiently **prepare your own** submissions.
Why?	✓ Speed ✓ Cost-Efficiency ✓ Simplicity	✓ Data Utility ✓ Defensibility ✓ Reputation	✓ Accuracy ✓ Volume ✓ Independence
Best if...	You'll need to publish a greater scope of documents under **EU CTR**, and you need to submit quickly and efficiently under EMA Policy 0070 and Health Canada PRCI.	You have clinical documents of public significance, and/or you want to honor the explicit preference of EMA and Health Canada for a **utility-preserving** approach.	You have a large, constant volume of submissions and an **in-house team** with experience handling clinical document submissions.

Why trust Privacy Analytics?

14 of the top 15 global pharma companies listed on PharmExec's Top 50 have trusted our solutions.

We have supported more than 100 successful regulatory submissions since 2017—with many more pending.

Every sponsor that has engaged with us in the last three years has re-engaged when they've had a subsequent submission.

Tight timelines? Need help?

Contact us to tailor an engagement model and a solution that meets your unique business needs.

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