

Protect Your Commercial Interest in Clinical Document Submissions



Customize a Solution to Protect Commercially Confidential Information (CCI)

Balancing the need for transparency with the need to protect CCI

With so much pressure to be transparent, how do you ensure your clinical study submissions don't reveal confidential trade secrets when published by regulators?

As a transparency leader, you must make clinical study documents public under regulations like EMA Policy 0070, EU CTR, and Health Canada PRCI while ensuring you don't disclose CCI. Most regulators accept only minimal redaction of these documents, requiring appropriate justification if a sponsor claims the data is CCI.

If you are...

- Preparing 1000+ page regulatory submissions in support of marketing authorization
- Concerned protecting CCI will disrupt how you redact or anonymize clinical documents for publication
- Unsure what qualifies as defensible CCI in your dossier or how to efficiently identify CCI at scale

...then Privacy Analytics' CCI Protection Services are for you.

Protect your competitive edge and brand trust with our CCI Protection Services



Competitive Edge

Protect your competitive edge with a sound strategy to identify and redact CCI in your documents, ensuring your trade secrets and IP remain private.



Peace of Mind

Protect your reputation while gaining peace of mind that you've safeguarded CCI *and* met transparency demands, influencing brand trust.



Speed at Scale

Safely accelerate clinical document submissions, on tight timelines, by identifying and redacting CCI and personal data efficiently and at scale.

Tailor a solution model to meet your needs

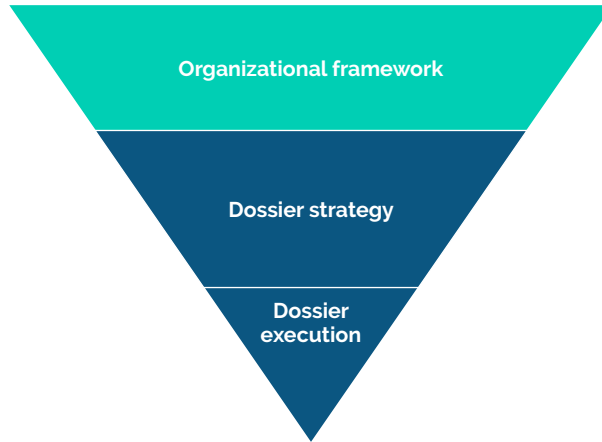
We offer a complete CCI protection solution – from strategy to execution – leaving you free to pick and choose the services and level of support required to meet your business needs.

“Low touch” support

We provide expert feedback on your CCI protection strategy.

You identify dossier-specific CCI, while we provide training and guidance to enable your team.

We provide templates with CCI justification options, enabling your team to redact CCI efficiently.



“High touch” support

We host workshops, creating a framework and guidance for your CCI protection strategy.

We consult, identify CCI, and run validation checks for your team's final review.

We redact CCI, incorporating justifications for regulators, which your team then approves.

Client success



According to a global pharmaceutical company in PharmExec's Top 10 that used our CCI Protection Services, success meant:

- ✓ Achieving internal alignment on what qualifies as CCI
- ✓ Defining a streamlined approach to redacting CCI
- ✓ Creating consistent justification tables that regulators approved

What could success look like to you? Set up a call to find out.

Why trust Privacy Analytics?

~95%

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

60+

We have supported more than 60 successful regulatory submissions since 2017—with many more pending.

100%

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

Need help protecting CCI? Let's talk.

Contact us to talk about a complete solution for your clinical document submissions.

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