redact360

Let's be honest. Manual redaction is complex, risky and expensive and striving to comply with new regulations takes up precious time you could be spending doing what you do best: finding cures for major diseases.

Thankfully, redaction and anonymization are what we do best.

Redact360 is an innovative new service from Kinapse that provides world-class, fully compliant redaction and anonymization supporting clinical trial transparency. At the heart of Redact360 is a sophisticated service: using an automated platform that ensures submissions meet rapidly expanding disclosure regulations (e.g. EMA Policy 0070, Health Canada PRCI), industry-driven transparency initiatives (EFPIA-PhRMA Principles for Responsible Clinical Data Sharing) and additional drivers (expanded journal requirements and public scrutiny).

~60 successful package submissions delivered and savings of over £1.5 million

With **Redact360**, you will not just fulfil your regulatory obligations:

- **You'll save time**
  By automating the identification and removal of direct and quasi-identifiers, our technology ensures that redaction timelines are reduced.
- **You'll save money**
  By outsourcing redaction and anonymization, you do not need to invest in technology or staff to deliver compliant submissions.
- **You'll drive efficiencies**
  You standardize all your submissions and channel them through a lean and hassle-free process.
- **You'll protect resources**
  You and your team are able to continue to focus on your main responsibilities.
- **You'll safeguard your reputation**
  Our 'continuous learning' environment draws on all our experience of previous submissions to make sure you become and remain an industry leader in compliant data sharing.

Redaction Process

Automation alone cannot deliver the quality required, and we are big believers in the power of human expertise.

Redact360 combines cutting-edge tech with a dedicated team of experts on hand to guide clients throughout the redaction process. Whether we are interpreting the constantly evolving EMA guidance, preparing 'marked for redaction' documents, pulling together the final redaction and anonymization package or helping you deal with questions that the EMA or Health Canada might have about your submission, you're supported at every step.

Finally, you can ensure compliance without stress. So how does it work?

We understand that every client is different. Your priorities and the ways you work are unique. We tailor our approach and process to make sure you get a personalized service that delivers exactly what you need, when you need it, in a way that feels effortless to you. We have global capabilities and we also understand that great communication is essential for time-constrained and highly pressured teams. We maintain a regular and well-structured dialogue with you throughout the entire **Redact360** process. You will receive insights and recommendations at every stage as your redacted packages are successfully submitted and your disclosure obligations are met.

For a free demo contact us at hello@kinapse.com

or call us on +44 (0)1276 713 800.

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