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LIFE SCIENCES

# Digital Privacy & Clinical Research

**Chaucer Life Sciences have been working with organizations involved in the control and processing of clinical data to support them with their GDPR compliance.**

We have a dedicated GDPR Life Science team made up of Data Privacy specialists and Life Science experts, who understand and advise on the implications of the new legislation on all aspects of Clinical Data Processing.

Combining our expertise in data integrity, clinical trials and data privacy we support life sciences organizations of all sizes with meeting the requirements of the GDPR, using the data privacy goals of each organisation as the starting point.

## We offer a mix of services, designed to provide support and expertise where needed. Challenges Chaucer has helped clients with include:

- Support clinical trials in seeking Ethics Committee approvals
- Auditing current methods of recording and managing consent
- Advising on safe and legal transfer of data outside of the EU
- Managing Subject Access Requests
- Pseudonymization/anonymization
- Definition and responsibilities of Controllers, Joint Controllers and Processors
- Managing Right to be Forgotten requests
- Operating studies without consent for research purposes – Article 89
- Ensuring Data Controllers understand increased documentation obligations

## DPO SERVICE

Outsourcing your DPO function to a specialist UK provider, reduces the pressures of a full-time hire and secures a higher level of expertise for your budget. This active role is fulfilled by data privacy experts who each bring over 30 years of experience in consulting and industry (CTO / CIO). Our practical advisors are independent, making sure you get the advice that's needed. General advice relating to cybersecurity, new technology or processes is also at hand.

**Flexible service to meet your changing needs:**

- DPIA Awareness
- Risk Management
- Policy Review
- Privacy Framework Management
- Breach Communications
- Incident Management
- Compliance Management
- Controls Management
- Best Practice

## EU REP

**We are helping American and Canadian Life Sciences organizations meet their responsibility under Article 27, by providing an EU Representative service.**

This allows your data subjects (EU based employers, contractors and volunteers) and data privacy regulators to correspond in their local language and time zone.


Functioning just like an extension of your business, the service includes a single source of truth database, and full assessment of relevant practices, and managing SARs. We also keep you up to date with any relevant legislative changes and court cases.

For more information, please contact Paul or Michael below:

Robert de Souza
Head of Data Privacy
+44 7714 916 511
robert.desouza@chaucer.com

Michael Whitworth
Head of Life Sciences
+44 7770 702092
rmichael.whitworth@chaucer.com