A GDPR Code of Conduct for clinical research service providers

CR.GDPR provides a recognised framework that incorporates both GDPR and ISO 27001 requirements to help sponsors understand how participating service providers meet the highest privacy and security standards needed within the clinical research domain.

Accelerate Study Start-Up and Reduce Audit Costs with CR.GDPR

Selecting clinical research service providers requires rigorous, time-consuming data privacy and information security assessments. CR.GDPR streamlines this process by combining GDPR and ISO 27001 requirements into a single, sector-specific framework and doing rigorous assessments of service providers against that framework.

Why CR.GDPR is a Commercial Advantage for Sponsors:

- **Faster Time-to-Market:** Shorten supplier onboarding and study start-up cycle times to get your trials operational faster.
- **Direct Cost Savings:** Drastically reduce your expenditure on vendor data protection and information security audits.
- **Instant Risk Mitigation:** Gain immediate assurance that your CROs meet the highest privacy and security standards across all 27 EU Member States.
- **Streamlined Procurement:** Standardise and simplify GDPR discussions, eliminating lengthy legal and compliance bottlenecks.

Protect Your Trial. Validate Your Vendors

Don't let compliance delay your science. When evaluating your next CRO or service provider, look for the CR.GDPR Compliance Mark on our public registry.

EUCROF
European CRO Federation

Ensure your partners meet the highest standards at crgdpr.org