A GDPR Code of Conduct for clinical research service providers

Win More Sponsors and Streamline Compliance with CR.GDPR

As a CRO or service provider, proving your data protection and information security posture to sponsors is a constant, resource-heavy bottleneck. CR.GDPR—the EUCROF GDPR Code of Conduct for Service Providers in Clinical Research—is the only sector-specific framework that officially validates your compliance across all 27 EU Member States.

Getting certified doesn't just check a compliance box—it gives you a distinct competitive edge in every RFP.

Why CR.GDPR is a Commercial Engine for Your CRO:

- Accelerate the Sales Cycle: Eliminate endless, repetitive data protection questionnaires. A CR.GDPR Compliance Mark gives sponsors instant assurance, reducing onboarding times and converting your sales faster.
- Win More Competitive Bids: Stand out in RFPs by holding a recognised, sector-specific privacy and security mark that competitors who don't adhere simply cannot match.
- Reduce Administrative Waste: Stop burning operational hours on conflicting, client-specific interpretations of the GDPR. Operate under one clear, universal standard, endorsed by the regulatory authorities in the EU.
- Gain Global Trust: Secure a tier-based Compliance Mark and a permanent slot on the official public registry, making your high standards visible to global sponsors and clinical research participants and stakeholders instantly.

Turn Compliance Into a Competitive Edge

Stop defending your data privacy posture on a client-by-client basis. Prove your excellence upfront.

EUCROF
European CRO Federation

Start your adherence assessment today at crgdpr.org