

Copernicus Group IRB



Major products/markets served

Copernicus Group IRB reviews studies for single investigators and serves as central IRB for multi-center trials involving several thousand investigators. Additionally, CGIRB functions as the IRB of record for studies conducted at hospitals and academic institutions.

Major services

Thoughtful ethical review provided by CGIRB is enhanced by the support of knowledgeable professional staff members. Copernicus Group IRB services include:

- Multiple IRB meetings each week
- Single Point of Contact
- 2-3 business day turnaround on documents
- Multi-center or single site reviews
- Education and Training Resources
- Dedicated Staff for institutional studies

Experienced in the Review of:

- FDA and OHRP Regulated Studies
- Phase I-IV studies in all therapeutic areas
- Device Studies
- Observational/Registry Studies
- Social-Behavioral Studies

Corporate description

Copernicus Group IRB was established in July 1996 as an independent institutional review board and operates in compliance with Federal Regulations governing IRBs set forth in 21 CFR and ICH GCP Guidelines. CGIRB is a member of the Consortium of Independent Review Boards (CIRB), which regularly meets with government regulators to review issues of importance to the protection of human subjects involved in clinical research.

In August 1999 and July 2005, Copernicus Group IRB underwent successful FDA inspections (no FDA Form 483 issued). In July 2004, CGIRB's Human Research Protection Program was awarded full accreditation by AAHRPP®, The Association for the Accreditation of Human Research Protection Programs, Inc. Copernicus Group IRB is a Certified Women's Business Enterprise and can provide a certificate to assist clients in meeting supplier diversity initiatives.



Copernicus Group IRB

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