Study Review Solutions

A single source of expertise for ethical and scientific review.

For more than 55 years, the industry has relied upon WCG as the gold standard provider for IRB services. We know how to navigate clinical trials. We have deep experience with virtually all sponsors, CROs, Institutions, and research sites in the industry – encompassing thousands of trials, across all phases and therapeutic areas. Clients tell us they respect our commitment to delivering with the highest quality, efficiency, and accuracy, in order to support timely decision making.

Today's clinical trial landscape includes more trial starts, growing protocol complexities, and more resource constraints. But one focus stays the same: The need to protect research participants through independent review of clinical trial safety and efficacy data. WCG offers clients the ability to leverage one expert organization for all your ethical and scientific review needs for IRB, IBC, DMC (or DSMB), and EAC (or CEC) review

EXCEPTIONAL EXPERTISE

With WCG Study Review Solutions, you have access to over 1,200 independent experts in our global network. Flexible, comprehensive services include a high level of expertise and efficiency:

IRB – Institutional Review Board

- Full board determination of a new protocol within 5-6 business days of a complete submission
- Site approvals within 1-2 days of complete site submission
- Review of site lists with insights on each institution
- Contact with key institutional players for IRB reliance questions
- Dedicated client-specific point of contact

IBC – Institutional Biosafety Committee

- Critical for development with engineered DNA or RNA as human gene transfer products
- Transparent and interactive, allowing sites to understand and direct compliance activities
- Bundles IRM and IBC review to eliminate duplicative reviews and ensure efficient turnaround

DMC – Data Monitoring Committee

- Our approach to DMCs place emphasis on quality of statistical analysis and committee experience to ensure our clients receive the most comprehensive and actionable recommendations to inform their pipeline development
- Proven team of statisticians with experience across over 25 therapeutic areas
- We understand the value of an effectively conducted DMC to protect patients, promote trust and support regulatory submissions

EAC - Endpoint Adjudication Committee

- Purpose-built, proprietary technology for streamlined Adjudication Committee management
- 5 days average time from posting to adjudicator decision
- 100% on-time final adjudication database lock (DBL) prior to final study DBL
- Critical for type II diabetes (cardiac adjudication), non-alcoholic steatohepatitis (NASH), and muscular dystrophy protocols

Contents

1,000+

DMC and EAC Expert Committee Members

200+

IRB and IBC Expert Committee Members

55+

years as the gold standard provider for Independent Review Services

The organization that invented the Study Review industry continues to reinvent it.

Experience is the key to fast, efficient, and accurate study review. Over 55 years' experience across all trial phases and therapeutic areas makes WCG the gold-standard for ethical, scientific, and regulatory review. Get the answers and insights you need with the industry's largest network of vetted sites and accredited committee experts.

Talk to our team today for a complimentary consultation

SCHEDULE A MEETING