



75+

FDA Approvals
Supported

50+

Years of Company Experience
Managing Committees

1,000+

Experts in Our Global Network

How Our Clinical and Statistical Expertise is Helping Sponsors

WCG's deeply experienced scientific leadership team combines more than 200 years of collective industry expertise in designing specialized clinical oversight and statistical solutions.

We help clients **drive efficiencies beyond traditional approaches** to achieve high-quality trusted data, exceeding regulatory requirements, and increasing the likelihood of approval.

Engage Independent Expert Endpoint Adjudication and Data Monitoring Committees in a strategic manner to identify methods for retrospective and prospective evaluations that may improve competitive advantage and bolster regulatory submissions.



Apply up-to-date knowledge of regulatory requirements and evolving committee best practices with industry-leading technologies to achieve process excellence and avoid unnecessary infrastructure spend.

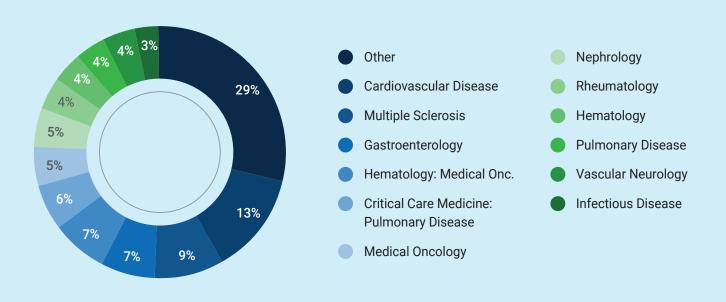
Understand and define risks early for timely informed decisions and course corrections that may **optimize trial spend** and commercial advantages.





Global Expert Network

WCG brings clients the largest committee dedicated staff in the industry, our own global network of more than 1,000 vetted medical, safety, and statistical experts across therapeutic areas who can serve as committee members or expert advisors to clients, and proprietary technology platforms that provide user-friendly data packages and streamlined operations for effective decision-making.



WCG's in-house independent statistical reporting group is comprised of over 40 biostatisticians and statistical programmers. Our biostatistical team's depth of experience and unique advantage in supporting DMCs stems from our years of experience on all sides of the table. Sponsors and DMC members widely recognize WCG for the accuracy and clarity of our monitoring reports and practice of gentle guidance.

We serve as scientific partners to our clients in trial design and regulatory strategy consulting, DMCs, statistical programming, and a variety of other services to help analyze your data and prepare your trial for regulatory submission and review.