

# Optimize Your Research Program to Speed Start-up, Support More Trials, Enroll More Patients, and Improve Financial Results

Integrated Research Administration Solutions Designed to Optimize Your Research Program Throughout the Entire Clinical Trial Lifecycle.

## Ethical Review & Biosafety

- IRB
- IBC
- IRBNet

## Study Start-Up

- Coverage Analysis
- Contract Redline
- Budget Development
- Contract Negotiations
- Budget Negotiations
- CTMS Study Builds

## Study Oversight & Management

- Billing & Receivables Management
- CRC & CRN Support
- DMCs & EACs
- Study Advertising
- Data Entry
- Statistical Consulting
- Patient Recruitment
- Patient Screening & Scheduling
- Imaging
- Velos eResearch CTMS
- Regulatory
- Florence eReg

## Study Close-Out

- Financial Audits
- Data Entry

## Ongoing Program Growth

- Sponsor Matching
- Industry Trainings & Resources
- Benchmarking
- Quality Consortium
- Consulting & Advisory
- WCG Site Network

## Benefits of Partnering with WCG



**Outsource administrative tasks** to optimize your research program throughout the entire clinical trial lifecycle and beyond.



**Leverage proven processes and technology** to accelerate study start-up and study management while ensuring quality and compliance.



**Alleviate resource constraints** and **reduce staff burden** so your existing teams can focus on patient care.



**Gain a trusted partner** that supports your site's exact needs, allowing you to build a more durable research program that grows with you.



**Reduce costs**, implement expert financial management processes, and build a more financially sustainable research program.



**Support more life-changing trials** and **expand the available care options** for your patients and community.

## Real Results

Increase Enrollment  
Rates by

**44%**

Reduce Study Start-up  
Timelines by

**37%**

Improve Financial  
Results by

**21%**