Westat®

Commercial Life Sciences

Moving Forward Together

Westat®

Improving Lives Through Research®

How can Westat help you?

SERVICES

Biostatistics

We provide comprehensive biostatistical support for multisite investigational studies in all phases of clinical research as well as expert study design and data analysis capabilities in collecting and interpreting appropriate study data.

CDISC Standards

Westat has fully implemented the Clinical Data Interchange Standards Consortium (CDISC) standards required for data management, programming, documentation, and eCTD submissions.

Clinical Trials Design

Our experts design efficient, rigorous clinical studies that ensure high probability of success, scientific acceptance, and regulatory requirements.

Data Management

Westat is trusted to develop and manage cost-effective, efficient, optimally designed data management systems that deliver robust data with rigor and transparency.

Data Monitoring Committee Support

We offer a full complement of services from staffing and drafting the charter to managing logistics. Our dedicated biostatisticians perform reliable, end-to-end reviews and analysis of safety and efficacy data to support your critical decisions.

Medical Writing

Westat's professional medical communicators develop accurate, complex publications and dossiers that are scientifically and ethically sound, compliant, and directed to the right target.

Pharmacovigilance

Our pharmacovigilance (PVG) team supports the product development lifecycle every step of the way—from early-phase studies to regulatory approval.

Project Management

Our project managers achieve project goals with an unwavering focus on quality control, milestones and deliverables, budgets, and client satisfaction as well as day-to-day oversight of all operations.

Regulatory Affairs & Operations

Westat interprets and relays regulatory requirements to help mitigate risks and prepare and publish complex, compliant dossiers—all while helping sponsors through the FDA review and approval process.

Site Management & Monitoring

From site selection and management to monitoring, site contract negotiation, and implementation, Westat provides a full suite of comprehensive site-related services critical to the success of your clinical trial.

About Westat CLS

Westat CLS is more than a contract research organization (CRO). We are an extension of your team—a trusted partner from first concept to final regulatory reporting.

Why choose Westat for your Clinical Trials?

Innovation

While our forward-thinking researchers see the world through data and clinical evidence lens, we also focus on the humans behind the numbers. Our CLS teams deliver cutting-edge expertise to design patient-centered, robust clinical studies.

Efficiency

Westat understands the budget and schedule pressures involved with bringing new treatments to market— and our comprehensive, efficient processes reflect our commitment to delivering value rapidly while ensuring quality.

Creativity

Our resourceful experts pivot easily to creatively solve problems and address challenges that may arise in the course of clinical trials and respond nimbly to changes in scientific or business direction when needed.

Integrity

With Westat, you can count on direct, reliable communication with experts at the top of their field, forming a trusted partnership to help you seamlessly navigate the clinical research process.

Leading Clinical Trials Experts

When you work with Westat, you get integrity, trust, and our best team—every time. You'll have a direct line to experts at the top of their field who are invested in partnering with you to drive rigorous studies and outcomes to help advance innovative treatments through early to late phases of clinical trials.

CONTACT

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