VERISTAT

Full-Service Clinical Trial Delivery

Advancing a novel therapy from clinical development to approval and commercialization is a complex, high-stakes process. Veristat's scientifically grounded experts do more than execute clinical trial plans—they think strategically. Our collective teams assess challenges, generate and evaluate data, plan across multiple scenarios, and carefully weigh trade-offs and risks, all while aligning short-term execution with long-term goals.

In the last 5 years

+110 Full-Service Projects

50% Biologics

>30% Rare Disease

Clinical Monitoring

From protocol development through study closeout, clinical monitoring plays a critical role in supporting efficient and high-quality project execution. A proactive, data-informed strategy is implemented early to anticipate and address potential issues, with continuous assessment and refinement throughout the study lifecycle. Veristat offers a flexible, customized monitoring model—ranging from comprehensive source data verification to targeted endpoint monitoring—tailored to client needs and aligned with global regulatory standards.

Site Management & Documentation (SM&D)

At Veristat, the Site Management & Development (SM&D) team prioritizes purposeful and transparent communication, fostering strong, lasting partnerships with clinical sites. This collaborative approach consistently drives site engagement and compliance across key areas—including regulatory start up and essential documentation, participant recruitment and retention, data quality, and adherence to timelines. In a competitive, fast-paced trial landscape where site resources are often stretched thin, Veristat's strategic site management ensures your study remains a priority.

Clinical Operations & Trial Delivery

Expertise to Design, Start-Up and Manage Clinical Trials to Successful Completion

Feasibility + **Patient Recruitment & Retention** + **Site Management** + **Clinical Monitoring** + **Project Management**

Full-Service Success Stories

1 From Challenge to Success: Expert Management of a Pivotal Pancreatic Cancer Study

Veristat supported a Phase II trial of a synthetic peptide for stage IV pancreatic cancer, providing comprehensive services across 20 US sites. Despite challenges with recruitment, EDC systems, documentation, and site communication, the team implemented effective solutions including enrollment boosters and systematic reviews. The primary endpoint was achieved with no safety concerns, prompting early study closure and continuation to Phase III with Veristat's partnership. **Learn more ›**

2 Collaborative Problem-Solving in Rare Gene Therapy Research

Veristat supported a multinational Phase II gene therapy trial for a rare lysosomal storage disorder by recruiting qualified global sites when initial recommendations failed, implementing precise scheduling for time-sensitive therapy administration, and delivering customized tracking systems when expanded metrics were requested. Despite the study being halted due to disappointing efficacy results, the partnership continued with Veristat managing post-transplantation requirements through a long-term follow-up study. **Learn more ›**

3 Strategic Management of Complex Metabolic Disease Treatment Logistics

Veristat supported a biotech company's Phase I/II gene therapy trial for a rare genetic lysosomal storage disorder by managing time-sensitive therapy administration requiring coordination between screening, stem cell collection, and inflexible manufacturing dates. They facilitated expedited data reviews between cohorts despite treatment delays. Through meticulous planning and innovative solutions, Veristat demonstrated exceptional flexibility, leading to continued collaboration on a long-term follow-up study. **Learn more ›**