



# VERISTAT

Scientific integrity. Client focus.



## Clinical Monitoring

### ABOUT VERISTAT

Veristat, Inc., is a full-service clinical research organization with demonstrated expertise in supporting clinical trials for pharmaceutical, biotechnology, and medical device companies. From trial design to final study reports, we offer complete services through strategic partnerships with our clients for an entire clinical program. With over 15 years of experience, Veristat provides flexible, innovative, and science-focused services customized to our clients needs.

### OUR SERVICES:

- Clinical Monitoring
- Data Management
- CDISC Standardization
- Biostatistics
- Medical Writing
- Project Management
- Regulatory Submissions
- Strategic Consulting

### WHY CHOOSE VERISTAT

There are several advantages to choosing Veristat.

- We are flexible and scalable to our clients' needs, and offer diverse therapeutic expertise, with a heavy focus on vaccine and oncology trials.
- In our Sponsor partnerships, we have proven to be adaptable and accommodating, which is reflected in our client satisfaction and retention.
- We are leaders in utilizing the latest technology platforms to assist Sponsors in the challenges of adhering to new industry standards.
- We are committed to providing quality services with the highest ethical and scientific standards that meet or exceed the expectations of our clients.

Veristat offers comprehensive clinical monitoring services from site initiation to closeout. Our clinical services are tailored to our clients' needs, complimenting product development plans and overarching corporate and quality management policies. All monitoring activities are conducted in accordance with the study protocol, standard operating procedures, ICH, GCP, and applicable FDA regulations.

### Flexibility

For small early phase studies through large multi-center studies, we provide strategic monitoring plans to support all clinical programs. Significant emphasis is placed on flexible and creative problem-solving approaches that highlight efficiency and streamlining of activities, thereby providing optimal solutions for our clients. Further, our highly experienced clinical staff operates within an integrated and collaborative environment focused on teamwork.

### Expertise

Our clinical monitoring services are provided by qualified, knowledgeable professionals with extensive clinical research and therapeutic area experience. Veristat's regional Clinical Research Associates (CRAs) have minimum of 5 years of industry experience. Sponsors always have the opportunity to review CVs and perform reference checks prior to approving monitoring personnel for their study. This approach ensures maximum coverage and management for each program, and provides our clients with a skilled and qualified therapeutically focused team.

### Clinical Monitoring Services:

- Site selection & feasibility
- Patient recruitment
- Site qualification & initiation
- Interim monitoring & site management
- Safety reporting
- Regulatory document management
- Remote monitoring via EDC

### Clinical Project Management Services:

- Site contract administration
- Supervision & training of CRAs
- Risk mitigation planning
- Continuous quality control
- Timeline & budget tracking
- Preparation & implementation of SOPs
- Investigator meeting coordination

For more information, call us today at (508) 429-7340 or visit our website at [www.veristat.com](http://www.veristat.com).