VERISTAT

The Science-First CRO™ and Consultancy

Advance Your Revolutionary Therapy's Development, Approval, and Commercialization

For 30 years, Veristat has helped pharmaceutical and biotech companies accelerate their clinical trials and commercialize their therapies by delivering Science-First solutions across the development lifecycle.

From overcoming unprecedented challenges in drug and biologic study design, to efficient patient recruitment and retention for an ultra-rare disease, to intricate clinical trial conduct and scientifically vetted regulatory submissions, our global team of experts is big enough to tackle any unknown while small enough to give you the attention you need to hit your milestones faster.

Our Impact – Improving Patient Lives

Since our inception, Veristat teams have prepared 90 marketing applications that received regulatory approval, including the first gene therapy approved in Europe.

Full Service Collaboration Across Entire Development Lifecycle

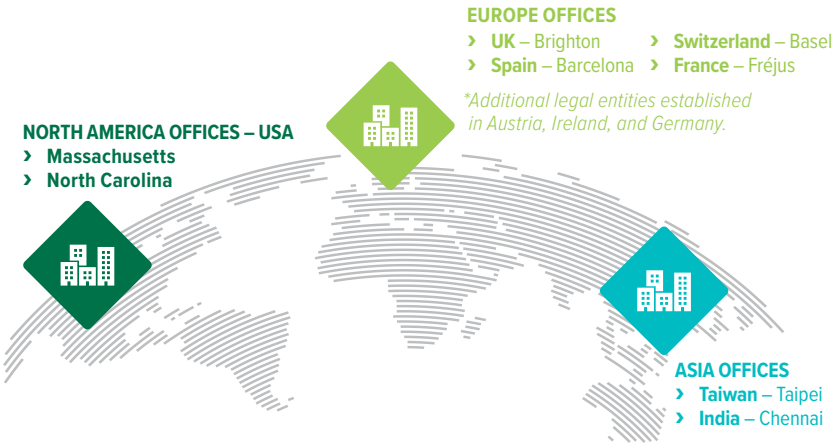
	PRE-CLINICAL	PHASE I-III	REGISTRATION	COMMERCIALIZATION
STRATEGIC CONSULTING	Scientific strategic insights to plan for development program success – from IND/CTA to post-approval			
REGULATORY AFFAIRS	Regulatory expertise that delivers the roadmap, gap analysis, agency interactions and registration/maintenance support to achieve success			
BIOMETRICS		Data and Statistical experts to plan the collection, analyses, standardization, and reporting of your clinical trial data		
CLINICAL OPERATIONS/ MEDICAL AFFAIRS		Integrated clinical trial conduct solutions including feasibility, site selection, patient recruitment & retention, site management, site monitoring, patient safety services and a flexible virtual/decentralized trials approach		
MEDICAL WRITING		Scientific-minded medical writers to develop protocols, clinical study reports, all modules of regulatory dossiers for global regulatory authorities, as well as scientific manuscripts to disseminate scientific and clinical data		
PHARMACOVIGILANCE			Safety and PVG specialists focused on managing the post-market safety risks of your approved medical therapies	
MARKET SOLUTIONS/ COMPLIANCE			Market Access, Reimbursement, Healthcare Compliance, Public Affairs and Quality Assurance expertise to ensure the commercial success of your product	

A Focus on Complex and Novel Therapies

In the past five years, Veristat has supported:

- **350+ rare/ultra-rare disease projects** and the preparation of **40+ marketing applications**
- **200+ projects** for endocrine/ metabolic disorder treatments
- **100 projects** for rare cancers
- **375+ projects** for **120+ sponsors** developing biologics
- **130+ projects** for cell and gene therapy

600+ Associates Across 3 Continents



"I would like to thank the entire team at Veristat, who worked day and night to submit our DSUR and our IND within thirty days of starting work with us! In addition to managing the process, the team worked seamlessly with us in the United States and our partner in Korea. Your team advised and guided us to produce quality submissions to meet our challenging corporate goal of submitting both on time. Due to these extraordinary efforts, we plan to work with you on our second IND submission."

CEO
Clinical stage biotechnology company

What You Can Expect from a Veristat Partnership

- Gains in quality, flexibility, and time
- Collaborative project planning and management
- Inter-disciplinary clinical, medical, statistical and regulatory guidance
- Flexible engagement models tailored to fit your program's specific requirements

Meet Veristat

The Science-First Full Service CRO and Consultancy That Delivers Results

If you haven't yet found the scientific expertise you had hoped for, with deadlines being missed and results falling short, talk to Veristat. We have a proven track record of starting—and keeping—clinical development programs on track with the flexibility to navigate the challenging path to regulatory submission and approval. Collaborate with experts who care like you do. It is not just business for Veristat, it's personal.

Meet Veristat Today | veristat.com