CORPORATE OVERVIEW FACT SHEET

VERISTAT

Accelerating Clinical Development and Regulatory Success

Delivering Speed, Expertise, and Results Across the Therapeutic Development Lifecycle

For 30 years, Veristat has helped pharmaceutical and biotech companies accelerate their clinical development programs and secure regulatory approvals by delivering comprehensive solutions that overcome complex challenges and meet critical milestones faster.

From navigating unprecedented challenges in study design to efficient patient recruitment in ultra-rare diseases to streamlining regulatory submissions, our global team combines deep expertise with personalized attention to accelerate your path to market.

Our Impact—Improving Patient Lives

105+ Marketing Applications that Veristat teams prepared have **received approval to date.**

Accelerating Development Across the Entire Lifecycle

	PRE-CLINICAL	PHASE I-III	REGISTRATION	COMMERCIALIZATION
STRATEGIC CONSULTING	Integrating science-based strategic insights to plan for development program success – from IND/CTA to post-approval			
REGULATORY AFFAIRS	Expedited regulatory expertise from strategy to execution and full registration support to achieve regulatory success			
BIOMETRICS		Data and statistical experts to optimize the collection, analysis, standardization, and reporting of your clinical trial data		
CLINICAL OPERATIONS		Focus on patient-centric clinical trial delivery solutions including feasibility, site selection, patient recruitment & retention, site management, and clinical monitoring		
MEDICAL WRITING		Scientifically precise medical writers to rapidly develop protocols, clinical study reports, all modules of regulatory dossiers for global authorities, and scientific manuscripts		
MEDICAL AFFAIRS SAFETY/PV			Patient safety and medical monitoring expertise throughout the lifecycle. PV specialists focused on managing the post-market safety risks of approved medical therapies	
MARKET ACCESS			Market Access, Reimbursement, Healthcare Compliance, Public Affairs, and Quality Assurance expertise to ensure rapid commercial success	

Proven Experience Across Novel Therapies

In the past five years, Veristat has accelerated:

- **480+ Oncology/Hematology** clinical trials and consulting projects
- **375+ projects for 120+ sponsors** developing biologics
- **350+ rare/ultra-rare disease projects** and the preparation of **40+ marketing applications**
- **200+ projects** for endocrine/metabolic disorder treatments
- **200+ projects** for neurology/psychiatry therapies

What You Can Expect from a Veristat Partnership

- Accelerated timelines with gains in quality and flexibility
- Proactive project planning and management
- Inter-disciplinary clinical, medical, statistical, and regulatory guidance
- Flexible engagement models tailored to fit your program's specific requirements and timeline objectives

400+ Associates Across 3 Continents

"Veristat provided strong data, very high-quality stats work, as well as fair, balanced, and compelling medical writing telling the story behind the data. Our sincere thanks for the partnership that made it possible."

SVP Regulatory Affairs and Quality Assurance

(NDA prepared by Veristat was approved in 2024)

Meet Veristat

Full-Service CRO and Consultancy That Accelerates Success

If you've struggled with missed deadlines and slow progress, talk to Veristat. We have a proven track record of starting—and accelerating—clinical development programs with the expertise to navigate the challenging path to regulatory submission and approval efficiently. Collaborate with experts who understand the value of speed without compromising quality.

It's not just business for Veristat, it's personal.

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