VERISTAT

Driving Successful Outcomes for Cardiovascular Trials

As leading life-shortening illnesses, cardiovascular diseases are critical to prevent and treat immediately in patients around the world. That is why Veristat has built a team of scientific-minded experts who have supported more than 120 cardiology and vascular projects, including drugs, biologics, and diagnostics. Our diverse trial experience, coupled with our knowledgeable and experienced teams, allows us to confidently handle the most complex trials and patient populations.

Our integrated global team has supported **>120 cardiology and vascular projects**

30% of cardiovascular programs are **rare/ultra-rare**

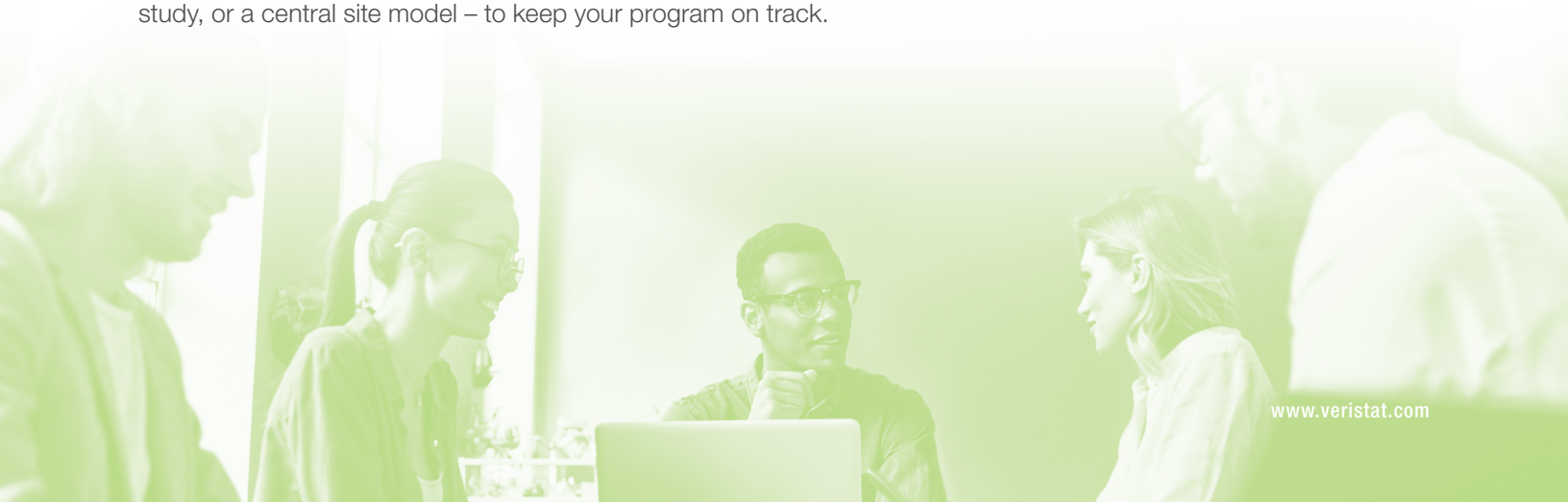
Specialized Expertise from Consultation to Beyond Submission

Pre-Clinical/Trial Planning to Increase Speed

Our diverse trial experience allows us to confidently handle the most complex trials and patient populations with the attention and focus needed to navigate the cardiology/vascular disease landscape. Veristat can determine if your study qualifies for an accelerated regulatory approval pathway and represent you in interactions with the US Food & Drug Administration (FDA), European Medicines Agency (EMA) and regional European regulatory bodies.

Study Design and Methodology that Will Work

Veristat's bold thinking ensures that your clinical trial or program design supports your regulatory strategy, whether you plan to run a single pivotal trial or multiple trials. We work with you to develop a customized solution, our goal being your success. We can help you select the right methodology – whether a decentralized trial, natural history study, or a central site model – to keep your program on track.



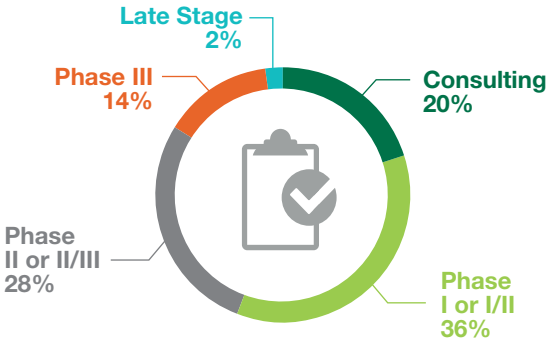
Marketing Application Preparation Experience by Phase and Publishing

When your next milestone is to get your NDA, BLA, MAA, NDS or jNDA submitted on time, trust our integrated global team that has prepared more than 160 marketing applications that have led to more than 80 approvals so far, with a 100% success rate in cardiovascular diseases.

Post-Market Pharmacovigilance Ensures Safety

Our post-marketing pharmacovigilance support team manages the safe use of your therapy and keeps it available to patients and their families.

Experience by Phase



Marketing Applications by Type

Bold Thinking that Delivers Results

We understand that all the easy challenges have been conquered in your clinical development programs and you still need to navigate the process and scientific unknowns. To ensure your success, select Veristat as your partner.

www.veristat.com