VERISTAT

Bold Thinking Overcomes Oncology Clinical Development Challenges

Scientifically minded to mitigate risks throughout oncology clinical trial design and execution

Supporting the development of cancer therapies accounts for **>30%** of Veristat's work

When the complexities of designing and executing cancer trials in a highly competitive and complicated market can mean delays in delivering urgently needed treatments to patients, it pays to have a trusted partner by your side. In the past five years, our experienced teams have helped clients with more than **300 oncology studies** and have worked on more than **30 US and European regulatory marketing applications** for cancer treatments, including many rare cancers.

In the past 5 years:

>300
Oncology Projects

>90
Rare Disease Projects

>30
Marketing Applications

Specialized Oncology CRO from Consultation to Beyond Submission

Pre-Clinical/Trial Planning to Increase Speed

We can help you navigate the complexities of designing and executing trials for both traditional cancer treatments and specialized therapies such as immunotherapies, vaccines, and cell and gene therapies. 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 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 offer a range of solutions – including decentralized trials, natural history studies, and a central site model – to keep your program on track.

Marketing Application Preparation and Publishing

When your next milestone is to get your NDA, BLA, MAA, NDS or jNDA submitted on-time, trust our integrated team that has prepared more than 150 marketing applications, including more than 30 oncology approvals.

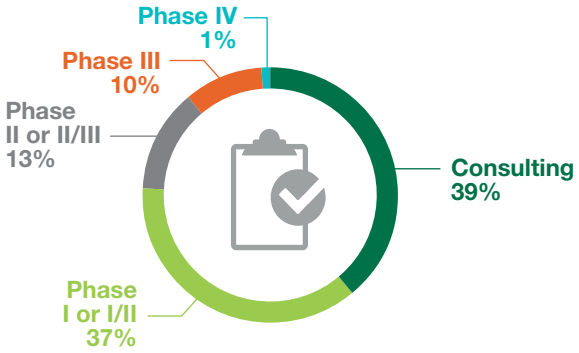
Post-Market Pharmacovigilance Ensures Safety

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

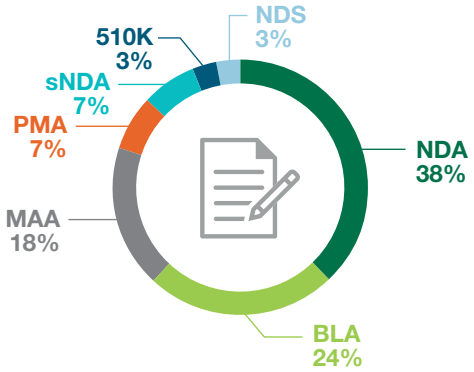
Oncology Expertise that Delivers Positive Impact

Veristat's teams have researched and worked on a spectrum of cancer treatments including drugs, vaccines, and diagnostics. Our efforts have brought about strong relationships and interactions with key global regulatory agencies, expertise in all expedited pathways to approval, and a strong record of success gaining approvals after one study.

Experience by Phase



Marketing Applications Prepared



*By partnering with a **like-minded, scientifically focused** company such as Veristat, we have found a **core team of high caliber collaborators** who can help us to accelerate the development of our new class of targeted cancer therapies. Our **preferred partnership agreement** with Veristat is a great step forward for our company given Veristat's **therapeutic and regulatory understanding** of both the fundamental elements of clinical research and the broader impact of our work"*

Chief Medical Officer, Mid-size Biotechnology Company

Contact Veristat Today

Learn more about Veristat and how we can assist you with your oncology trial development and execution.

www.veristat.com/therapeutic-expertise/oncology