CASE STUDY

VERISTAT

Rescuing Site Monitoring Success

Ensuring Continuity and Timely Completion Through Teamwork in Europe

Background

A European-based clinical stage firm focused on developing innovative immuno-oncology therapies engaged Veristat to provide succession monitoring due to their contract CRA retiring. While the study's first two cohorts were being run in 2 countries, Veristat took over the site management and monitoring activities for the final cohort, which included 3 sites in the United Kingdom.

The project team for the study was very small and included many functions provided by a multitude of different vendors. For example, the Project Management was being conducted by another CRO and they provided oversight of the Veristat CRA. Because all team members worked closely together in collaboration, the project successfully met the client's milestones.

Solution

Veristat joined the study team by assuming the responsibilities of the CRA who retired, which included monitoring the UK sites. To achieve the aggressive timelines, the Veristat CRA met with site teams on a weekly basis to ensure that recruitment, patient identification, and screening activity were on track. This continued until the recruitment of the needed 13 patients was complete.

In addition, the Veristat CRA worked meticulously with site teams to ensure the timely entry of data, allowing for the timely performance of data verification and database lock for each of the planned interim analyses.

STUDY DEMOGRAPHICS

INDICATION
- Pancreatic Cancer Vaccine/Immuno-Therapy

STUDY PHASE
- I/II

SITES
- 3 UK

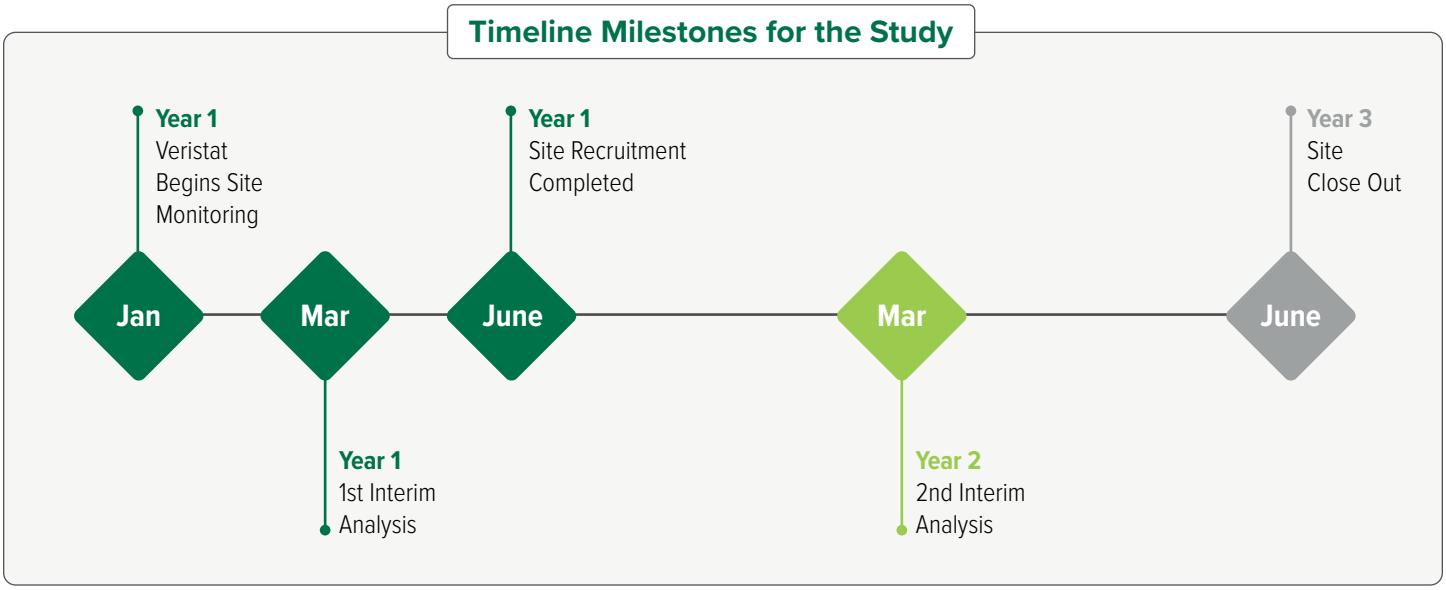
PATIENTS
- 13

SERVICES PROVIDED
- Site Monitoring
- Study Demographics

PROJECT TEAM
- 2 CROs
- 2 Central Labs
- IP Supply
- CRF Vendor

Goal

The sponsor required that first and second interim analyses be reached over the course of one year.



Result

Veristat completed the enrollment of 13 patients across the 3 UK sites on time. All data for the UK sites was entered and verified prior to the interim analyses. Veristat continued its interim monitoring visits for the sites and successfully closed the sites at study conclusion as planned and on time.

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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