PHARPOINT RESEARCH

EXPERTISE & SUCCESS STORIES

Supporting Oncology Clinical Trials

PharPoint's team has successfully participated in over 200 oncology trials since 2007 and has the right experience to seamlessly support your study.

Our team of experts have extensive experience within oncology, including experience with immuno-oncology, solid tumors, and hematologic cancers. This includes expertise within RECIST criteria and the STING pathway. We have experience with the FDA's oncology division since 2006, including a recent NDA submission for Chemotherapy-Induced Myelosuppression (CIM) and a BLA for NMIBC. Additionally, PharPoint's on-staff MDs have supported numerous oncology studies in their careers.

PharPoint's teams work collaboratively from the start to pursue a seamless study experience for sponsors, sites, and patients.

Keys to Success

Site Engagement Strategy

Industry-Best Timelines

SUCCESS STORY
Meeting Enrollment Goals Ahead of Schedule

In one recent oncology success story, PharPoint partnered with a sponsor to provide project management, clinical monitoring, site management and biometrics support.

The study, a Phase 1 trial within breast cancer, enrolled 17 subjects in 4 escalating-dose cohorts across 4 clinical sites. PharPoint's team facilitated enrollment adjudication in partnership with the sponsor.

Each cohort filled within the expected time frame, and Last Patient Last Visit (LPLV) was completed ahead of schedule.

SUCCESS STORY

Oncology Rescue During COVID-19 Pandemic

During the COVID-19 pandemic, PharPoint was brought onto an oncology study as a rescue partner.

Our team pivoted the study to remote monitoring,adjusting monitoring schedules per site capabilities. Additionally, we guided sites through COVID-19 vaccine implementation in conjunction with the dosing schedule.

With support from our team, the study successfully sustained its enrollment rate.

Keys to Success

- Flexibility
- Consultative Approach
- Remote Monitoring

OUR EXPERT ADVICE

Areas of Focus for a Successful Oncology Study

- **Site Selection:** Leverage site expertise to maximize patient enrollment and engagement
- **Site/Patient Centricity:** Minimize site and patient burden to maximize engagement
- **Data Entry:** Stress importance of timely data entry during site qualification visits with the goal of minimizing backlog. If lengthy data entry processes are noted at a site during qualification, consider pursuing alternative locations
- **Investigator Budgets:** Recognize inflationary impact on investigator budgets during negotiations to allow for rapid CTA and site activation
- **Patient Profiles:** Ensure clinical collaboration with data management team to increase efficiencies in both monitoring and data cleaning
- **Patient/Cohort Management:** Ensure study team has tools to enable real-time review of patient data, allowing team to recognize critical signals as soon as possible

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ABOUT

PharPoint Research is an award-winning and client-focused contract research organization (CRO) that offers Phase 1 through Phase 4 clinical trial support to clients of all sizes. Founded in 2007, PharPoint's team has participated in over 1,000 clinical trials with a high business repeat rate.

Our services include:

- Clinical Operations
- Medical Monitoring
- Project Management
- Data Management
- Biostatistics
- Statistical Programming
- DMC Support
- Medical Writing
- Study Rescue