PHARPOINT RESEARCH

EXPERIENCE OVERVIEW

Supporting Immuno-Oncology Trials

PharPoint Research, Inc.

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About PharPoint

PharPoint Research is an award-winning and client-focused contract research organization (CRO) that offers site feasibility, clinical operations, data management, biostatistics, and statistical programming services 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.

By the Numbers

Our experts have supported numerous oncology and immunotherapy studies throughout their careers, including experience with treatment modalities and safety event adjudication.

200+ oncology trials supported since 2007

30+ immunotherapy studies supported

50+ oncology indications supported

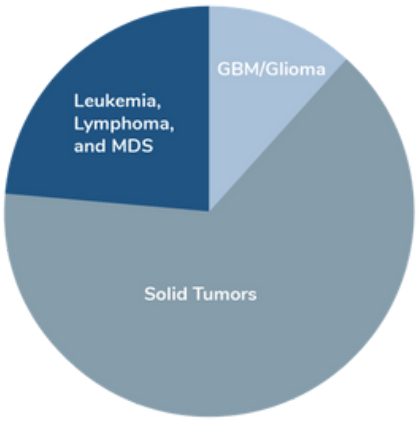
10 successful Oncology rescue studies

2 successful oncology regulatory submissions in the past 3 years

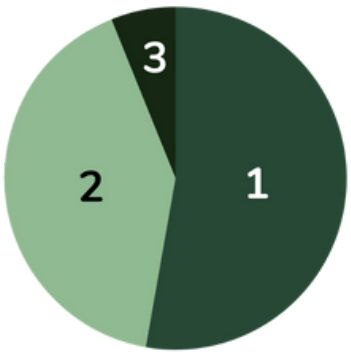
Industry-leading biometrics timelines, allowing you to get results **31 days faster** than industry average

Immuno-Oncology Breakdown

BY INDICATION:



BY PHASE:



Experience supporting studies with immunotherapy types including:

- Monoclonal antibodies and immune checkpoint inhibitors, including STING with PD-1/PD-L1
- Non-specific immunotherapies
- T-Cell therapies, including CAR-T and BiTE
- Oncolytic and Virus Vector Therapy

We have a strong appreciation for:

The need for **constant vigilance for safety monitoring and protocol adherence**, particularly regarding to immuno-oncologic therapy related safety events

Prompt and frequent communications with all stakeholders, including the need to establish trusting relationships with sites

The critical need to **recognize, manage and communicate adverse events**, and adjust the study interventions per protocol with the Sponsor's involvement

The high value of **leveraging a Sponsor's science, pre-clinical, and clinical experience** in decision-making during a study

Equipped to overcome common immuno-oncology study challenges:

CHALLENGE 1

Identifying and engaging the right sites

PharPoint's feasibility team identifies sites with access to patients & experience managing both patient care and logistics. We leverage our site relationships and knowledge, including an understanding of administrative lead time, to expedite FPI & keep your study top-of-mind.

CHALLENGE 2

Mitigating logistical risks associated with trial materials

Immuno-oncology trials are logistically complex, and our team plans ahead to mitigate risk and allow for a smoother study experience for all stakeholders. Our oncology-experienced PMs have expertise in managing logistics at a subject, site, and study level.

CHALLENGE 3

Managing increased safety complexities

Patient safety is at the center of all clinical trials. PharPoint's clinical team understands common adverse events and clinical indicators, and CRAs have experience and training using CTCAE. Our team also has experience managing studies through various protocol amendments.

Learn more about how PharPoint can support your upcoming study.

PharPoint Research is an award-winning and client-focused contract research organization (CRO) that offers clinical operations, data management, biostatistics, statistical programming, and strategic clinical trial consulting services to clients of all sizes.

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Speak with our team: pharpoint.com/contact