



Your experienced early phase development team.

PharPoint Research, an award-winning CRO, and **High Point Clinical Trials Center**, the largest clinical research facility in North Carolina, are partnered to provide sponsors with seamless Phase 1 study design and conduct experience.

Faster Timelines

With a robust recruitment database, dedicated site staff, and industry best timelines for support services, our teams ensure timely results.

Time-Cost Savings

Due to our long-standing partnership and geographical proximity, our repeatable proven processes create cost savings that we pass along to our clients.

Client-Focused Understanding

With a detailed understanding of early phase development challenges, we align study execution strategies to your needs and objectives. 70% of PharPoint's project managers have worked Sponsor-side, & High Point CTC was founded as an internal research arm of a biotech company.

70%

of studies conducted at High Point Clinical Trials Center are Phase 1

41%

of studies supported by PharPoint are Phase 1



PharPoint Research AWARD-WINNING CRO

- **FIH expertise:** Experience supporting studies evaluating pharmacokinetics & safety in normal healthy subjects
- **High client retention rate:** Services include clinical operations, project management, data management, biostatistics, and consulting
- **The right team:** Award-winning teams have relevant experience including supporting SAD/MAD, and adaptive trials
- **Get results faster:** Industry-best biometric timelines (one month faster than the industry average)

High Point CTC LEADING PHASE 1 UNIT

- **Purpose-built:** Large configurable space with dedicated staff, stand-alone pharmacy and specimen preparation laboratories.
- **Participant access:** 8,500+ healthy volunteer database
- **Proven recruitment:** Study-specific strategies developed by in-house recruitment team
- **Participant retention:** 98% participant retention rate
- **24/7 Oversight:** Fully trained and dedicated staff