CAPABILITIES

Global Service Solutions

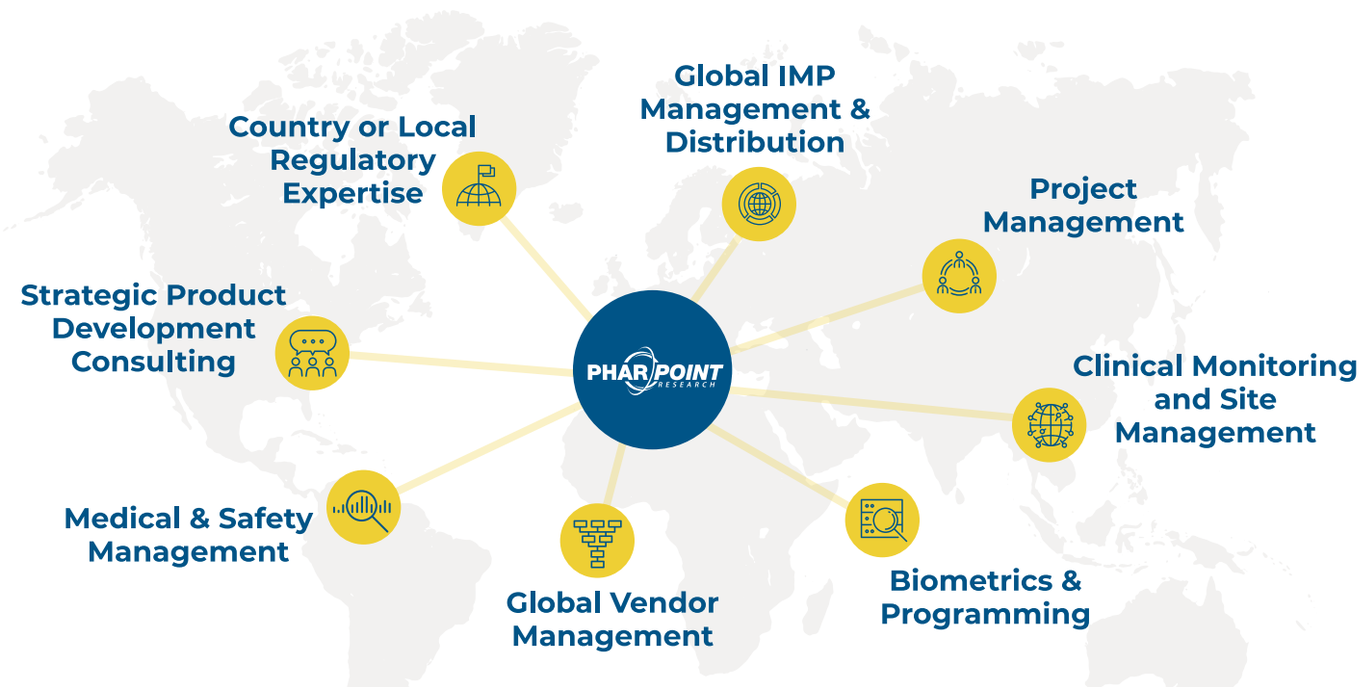
Through global partnerships, PharPoint's fully integrated, multi-disciplinary team is tailored to meet the needs of your study, equipped with consistent communication and a full knowledge of study direction.

For global studies, it is imperative that the study team have as much country and regional specific knowledge and resources to navigate the complicated regulatory environment unique to studies in multiple countries. PharPoint has vetted global CROs in Europe, Latin America, and Asia Pacific to allow us to choose the right partner based on study needs.

These resources would provide the regulatory and country specific knowledge needed but would report into the same Project Management structure as the North American based monitoring team to ensure consistent monitoring and clinical trial management.

PharPoint will work with our global partners to provide strategic planning and regional expertise for recommendation for country mix and navigation of regional regulatory requirements. PharPoint will coordinate the global team's efforts to optimize regulatory approvals, ethic committee submissions/approvals, enrollment, and proactively manage obstacles such as subject screen challenges and excessive dropouts at the study, region, and site level.

We look forward to working with our clients to find flexible solutions to meet their study needs.



Work with our award-winning team.

- **2020** *Triangle Business Journal Life Sciences Award Winner*
- **2019** *Scrip Awards Best CRO - Specialist Providers Finalists*

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