CLINICAL TRIAL CONSULTING

Guidance and consultative support that allows your team to successfully plan and design a clinical development program

PharPoint's network of regulatory, clinical, medical, and biometrics consultants provide high-level strategic support to help guide your team from pre-IND to regulatory approval. Equipped with our extensive knowledge and foresight to overcome study challenges, you can confidently make informed decisions and advance your product along the development pathway.

Program Strategy

Including:

- IND submission support/module development
- Strategic development planning and maintenance
- Program review support

Regulatory Strategy

Including:

- Regulatory strategy development
- Expedited pathway exploration
- Regulatory meeting support and representation

Trial Planning and Conduct

Including:

- Protocol design
- Phase 1-4 clinical trial planning and risk mitigation
- Key Opinion Leader (KOL) network development
- Investigative site feasibility

PharPoint Research has supported over 1,000 clinical trials. *To learn more about how we can support your current or upcoming clinical development program, meet with our team.*