SITE FEASIBILITY SUPPORT

PharPoint allows you to strategically identify and activate sites that fit your study needs.

Great sites give your trial a better chance at success. PharPoint's objective analysis and insight can help you effortlessly identify effective sites that can help you meet study goals.

Our robust feasibility process evaluates data including:

- **Patient availability:** Including looking at medical claims
- **Site performance:** Evaluating historical data on experience/enrollment in similar studies
- **Activation timelines:** For example, an evaluation of a site's ability to work within the Accelerated Clinical Trial Agreement (ACTA) framework and central vs. local IRB usage

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FEASIBILITY SUPPORT SERVICES MAY INCLUDE:

Investigator & Site Identification:
PharPoint can perform a data-driven site search to identify viable site options on your behalf.

Feasibility Assessment & Recommendation:
Let our experts assess identified sites to determine a best-fit for your study needs, and receive a tiered recommendation for sequencing qualification, initiation, and activation.

Essential Document Management:
Integrate our resources into your own and let our experts handle study startup details to support the regulatory readiness of sites.

Identify and activate sites faster

INDUSTRY FACTS

1 in 4 Sites

REPORT START-UP TIMELINES OF

121+ Days*

**Measured from Protocol Delivery to Site Activation, according to data from WCG Knowledge Base*

PHARPOINT

MANAGES RELATIONSHIPS WITH SITES WHO ARE ABLE TO ACTIVATE

Within 60 Days