SUCCESS STORY

Supporting a Client Who Met Study Enrollment Goals 5 Months Ahead of Schedule

ABOUT THE CLIENT:

A pharmaceutical company looking to conduct a Phase II dermatology trial in the U.S.

PHARPOINT'S ROLE:

PharPoint provided clinical operations, data management, and biostatistics support.

An innovative pharmaceutical company evaluating a new treatment option for a chronic skin condition chose PharPoint Research to provide clinical and biometrics support for their Phase II trial.

Due to equipment scarcity and timeline constraints, only two U.S. sites were eligible to support the study.

Of these two sites, one had **limited clinical research experience.**

While utilizing research-naive sites for a study requires added time and resources to ensure site staff are up-to-date with current guidelines and expectations, the study team felt both sites were needed to meet enrollment goals.

As both sites were activated, PharPoint and our client ensured excellent support and realistic goals were set for the site that was newer to research.

When the enrollment period began, a second, major unexpected challenge arose: **the COVID-19 pandemic.**

The study was placed on pause as site closures occurred across the U.S. In an effort to gain back time lost during this closure, a specialized patient recruitment partner was brought onto the study team.

Once the study's two investigative sites reopened and the study was resumed, they were flooded with prospective patients.

As subjects were enrolled, PharPoint's team maintained a collaborative relationship with both PIs, providing front row seats to our client regarding study progress. This included weekly updates for projected subject visits as well as closely monitoring and responding to missed visits.

The study was successfully, fully enrolled five months ahead of schedule.