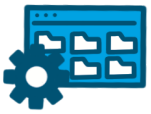
SERVICE OVERVIEW

Data Management and Biostatistics Clinical Trial Support

Thanks to experience, efficient processes, and excellent communication, PharPoint's U.S.-based biometrics teams help clients expedite timelines and meet study goals.



Clinical Data Management

Partner with a data management team that prioritizes your data from day one.

INDUSTRY-BEST TIMELINES

30 day
custom database builds, compared to an industry average of 68 days

SERVICES INCLUDE:

- Database build
- Data quality documentation
- CRF completion guidelines
- Across study data standardization
- SAE and external data reconciliation
- Medical term coding
- Consulting and oversight

Biostatistics and Statistical Programming

PharPoint's consultative approach to our biostatistics services includes a level of responsiveness, support, proactiveness that is unmatched within the industry.

NUMBERS AT A GLANCE

30+
regulatory submissions supported

100%
of statisticians have advanced degrees

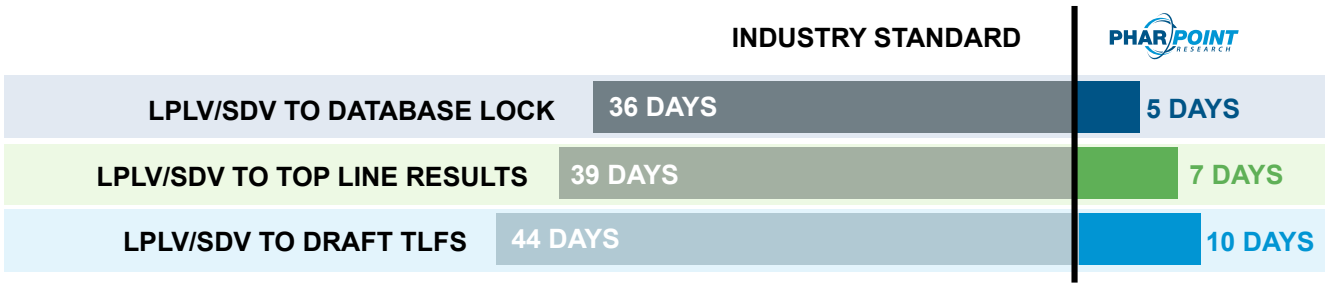
SERVICES INCLUDE:

- Randomization
- Regulatory and advisory meeting support
- Statistical analysis plans (SAP)
- Statistical analysis and support services for interim, futility, and DMC analysis
- Integrated summaries of safety and efficacy
- Preparation of case report tabulations (CRTs)
- Quantitative epidemiology including patient registries

GET RESULTS FASTER

with PharPoint's Industry-Best Timelines

At every step, PharPoint's standard biometrics timelines are industry-best. Throughout your study, clean data are soft locked, reducing the number of subjects needing to be locked following Last Patient, Last Visit (LPLV) and decreasing the timeline to database lock to only 5 days after final Source Document Verification (SDV).



AVOID HIDDEN COSTS

with Custom Programming

PharPoint's statisticians use custom programming, transferring ownership to clients at the end of a study **without any added costs.** Other CROs often use proprietary macros for their programs, which can lead to problems for sponsors during the submission process.

SAVE TIME AND MONEY

with PharPoint's Technology Partnerships

The PharPoint team has held partnerships with Medidata and Medrio since 2011, including in-house certified study builders and administrators. These partnerships allow us to pass on cost and timeline advantages to our clients.

CONFIDENTLY NAVIGATE REGULATORY DISCUSSIONS

with An Experienced Biometrics Team

PharPoint biostatisticians can support discussion with regulatory authorities, with experience providing statistical representation for Type A, B, and C meetings.