
# Rapid Success!

*PharPoint's Formula for Delivering Speedy Results*

**Time is money!** We have all heard this age old expression, but PharPoint Research takes this message to heart. Time is often a client's most limited resource, which is why PharPoint works diligently to deliver results faster than our competitors. Here's how:

**Flexible** PharPoint's flexible management structure is devoid of unnecessary layers of bureaucracy. Listening to the client's continually changing needs, our core executives are able to promptly make top level decisions, providing efficient solutions and cost savings.

**Consistent** PharPoint's consistent procedures reduce unwanted and unnecessary variation in data, allowing for the efficient use of resources and reduction of life cycle costs.

**Proactive** PharPoint listens to our client's needs in order to deliver personalized clinical trial solutions that anticipate the challenges of development. PharPoint's holistic and proactive management of each clinical trial exceeds our client's expectations by eliminating the headaches associated with the one-size-fits-all approach employed by larger CROs.

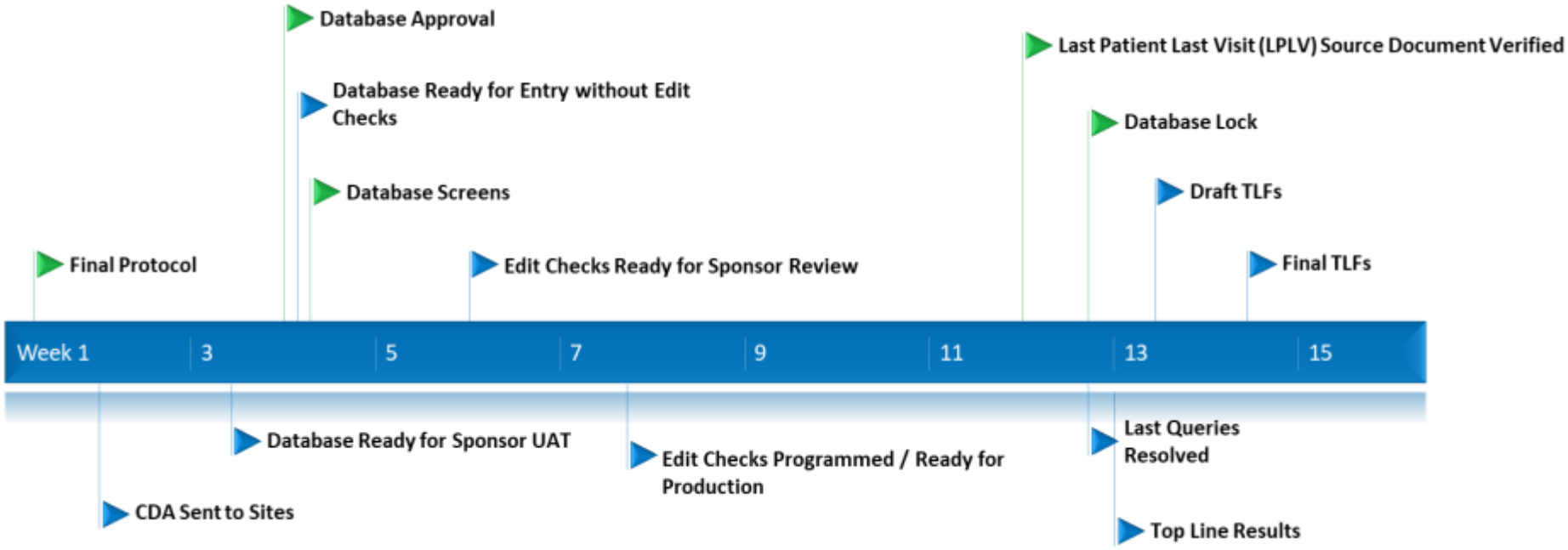

# Swift Timelines

What are these qualities important?

In essence, these qualities make up the foundation that allows PharPoint to deliver the following services in a fraction of the time it takes our competitors.

| Task/Milestone | Approximate Duration |
|---|---|
| CDA Sent to Sites | 1 Day after Kick Off Meeting |
| Site Recruitment Questionnaire (SRQ) | Within 1 Day of Executed Site CDA |
| First Site Qualification Visit | Within 5 Days of Satisfactory SRQ |
| Site Regulatory and Contract Activities | Within 1 Day of Sponsor Site Approval |
| Database Ready for Sponsor UAT | 15 Days after Final Protocol Received |
| Database Ready for Entry | 1 Day after Database Approval |
| Edit Checks Specifications Ready for Sponsor Review | 12 Days after Database Screen Approval |
| Edit Checks Ready for Production | 12 Days after Edit Check Specification Approval |
| First Site Initiated | Upon IRB Approval, CTA Budget Executed |
| First Interim Monitoring Visit | Within 2 Weeks of First Subject Visit |
| Last Queries Resolved | 5 Days after Last Patient Last Visit (LPLV) |
| Database Lock | 5 Days after LPLV has been Source Document Verified |
| Top Line Results | 2 Days after Database Lock |
| Draft TLFs | 5 Days after Database Lock |
| Final TLFs | 5 Days after receiving Draft TLF comments |

# Visual Representation




## Summary

PharPoint Research's flexible staffing structure is devoid of unnecessary costs or layers of bureaucracy. Listening to the client's continually shifting needs, our team delivers quality results in an efficient timeframe. PharPoint Research is ready to work with you to develop a strategy that meets your specifications and quickly gets your clinical trial on the path to success.