

# 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.

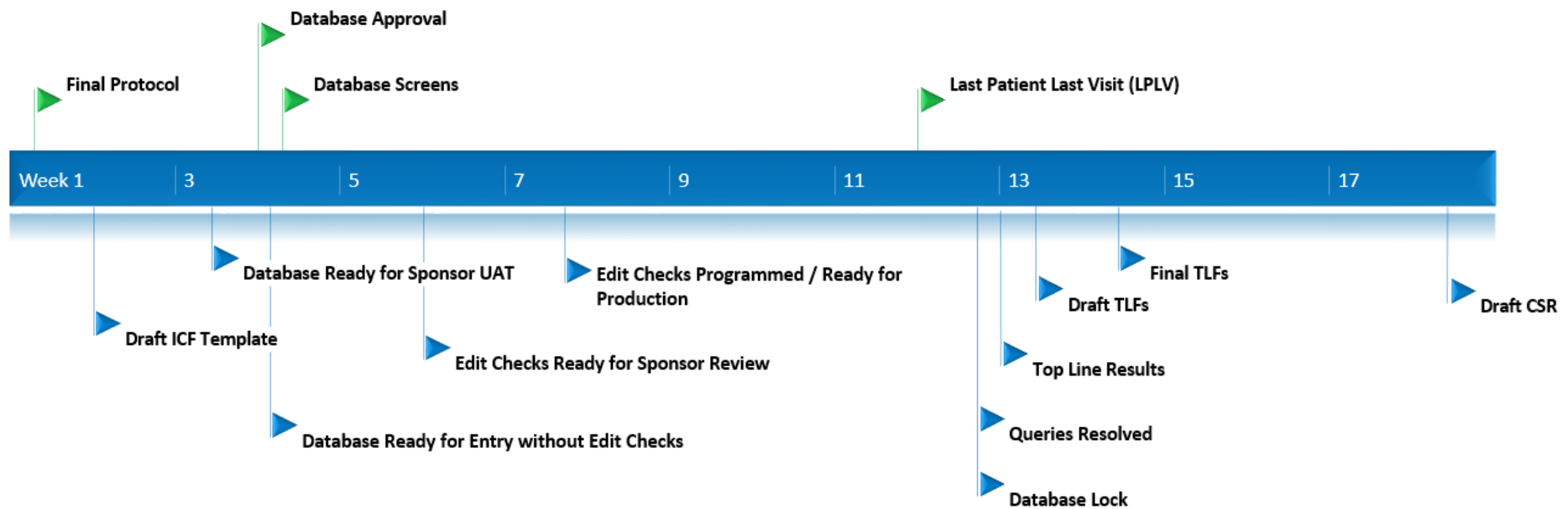
## Swift Timelines

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
Draft ICF Template	5 Days after Final Protocol
Database Ready for Sponsor UAT	15 Days after Final Protocol
eTMF Set up	1 Day after receiving Final Investigator List
Database Ready for Entry	1 Day after Database Approval
Edit Checks Ready for Sponsor Review	12 Days after Database Screen Approval
Edit Checks Ready for Production	12 Days after Edit Check Approval
First Interim Monitoring Visit	Within 2 Weeks of First Subject Visit
Queries Resolved	5 Days after LPLV
Database Lock	5 Days after LPLV
Top Line Results	2 Days after Database Lock
Draft TLFs	5 Days after Database Lock
Final TLFs	5 Days after receiving Draft TFL comments
Draft CSR	4 weeks after Final TFLs

## Visual Representation





## Summary

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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.