PHARPOINT RESEARCH

INDUSTRY-BEST

Standard Timelines

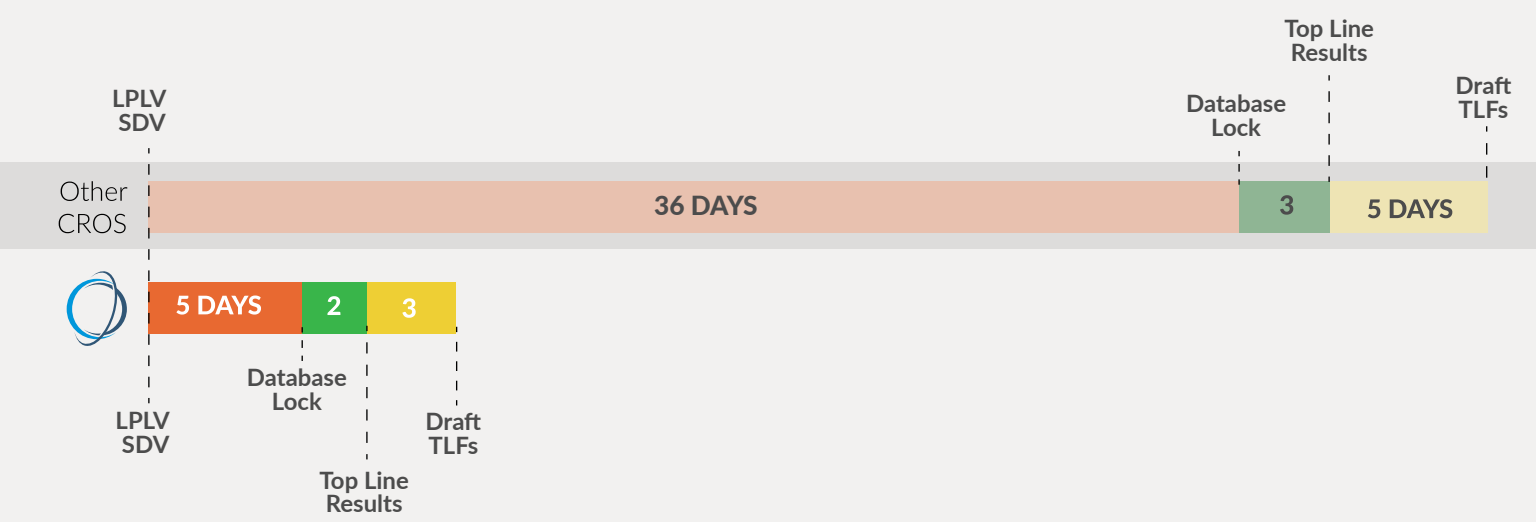
PharPoint offers clients industry-best timelines achieved through proactive planning and consistent cross-functional communication.

For information regarding expediting your upcoming study, reach out to a representative on our team at **bizdev@pharpoint.com**.

Task/Milestone	Approx. Duration
Database Screens Ready for Sponsor Review	**15 Days** after Final Protocol Received
Database Ready for Sponsor UAT	**5 Days** after Database Screen Approval
Edit Checks Ready for Production	**12 Days** after Edit Check Specification Approval
Last Queries Resolved	**5 Days** after Last Patient, Last Visit (LPLV)
Database Lock	**5 Days** after LPLV Source Data 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

Other CROs *vs.* PharPoint

To visualize the real difference between the **timelines of other CROs**** vs. the **PharPoint Standard Timeline**, we dove into the task-by-task details from Last Patient, Last Visit to the delivery of Draft TLFs.



** Dependent on country regulations & requirements*

*** Data used for "Other CROs" was collected from publically available information advertised by competitors as well as industry standards according to the 2017 Tufts CSDD industry report. Competitor timelines may vary.*

To set up a meeting, email us at **bizdev@pharpoint.com** or call **(919) 265-3655**